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JOURNAL      Submitted (26-JUL-1999) CBCB, Royal Ontario Museum, 100 Queen's
              Park, Toronto, Ontario M5S 2C6, Canada

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SOURCE      Amphibia: Batrachia: Anura: Neobatrachia: Bufonidae;
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REFERENCE   1 (bases 1 to 100)
AUTHORS    Liu,W., Lathrop,A., Fu,J., Yang,D. and Murphy,R.W.
TITLE       Phylogeny of East Asian bufonids inferred from mitochondrial DNA
             sequences (Anura: Amphibia)
JOURNAL     Mol. Phylogenet. Evol. 14 (3), 423-435 (2000)
MEDLINE    20179527
PUBMED     10712847
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REFERENCES  Liu,W., Lathrop,A., Fu,J. and Murphy,R.W.
DIRECT SUBMISSION Submitted (26-JUN-1999) CIOCB, Royal Ontario Museum, 100 Queen's
              Park, Toronto, Ontario M5S 2C6, Canada

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mrna
cds
features
source
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ACCESSION AF174512		
VERSION AF174512.1 GI:7620461		
KEYWORDS	Bufo melanostictus.	
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REFERENCE	1 (bases 1 to 100)	
AUTHORS Liu,W., Lathrop,A., Fu,J., Yang,D. and Murphy,R.W.		
TITLE Phylogeny of East Asian batrachians inferred from mitochondrial DNA sequences (Anura: Amphibia)		
JOURNAL Mol. Phylongenet. Evol. 14 (3), 423-435 (2000)		
MEDLINE 20179527		
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REFERENCE	2 (bases 1 to 100)	
AUTHORS Liu,W., Lathrop,A., Fu,J. and Murphy,R.W.		
TITLE Direct Submission		
JOURNAL Submitted (26-JUL-1999) CIBC, Royal Ontario Museum, 100 Queen's Park, Toronto, Ontario M5S 2C6, Canada		
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DEFINITION Bufo melanostictus KIZ-97J202 cytochrome b gene, partial cds;		
ACCESSION AF174513		
VERSION AF174513.1 GI:7620463		

KEYWORDS  
SOURCE  
ORGANISM  
REFERENCE  
AUTHORS  
TITLE  
JOURNAL  
MEDLINE  
PUBMED  
REFERENCE  
AUTHORS  
TITLE  
JOURNAL  
FEATURES  
source  
CDS  
mrna  
BASE COUNT  
ORIGIN

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Mitochondrion Bufo melanoscticus  
Eukaryota; Metazoa; Chordata; Vertebrata; Euteleostomi;  
Amphibia; Batrachia; Anura; Neobatrachia; Bufonidae;  
Bufo.  
1 (bases 1 to 100)  
Liu, W., Lathrop, A., Fu, J., Yang, D. and Murphy, R.W.  
Phylogeny of East Asian bufonids inferred from mitochondrial DNA  
sequences (Anura: Amphibia)  
Mol. Phylogenet. Evol. 14 (3), 423-435 (2000)

2 (bases 1 to 100)  
Liu, W., Lathrop, A., Fu, J. and Murphy, R.W.  
Direct Submission  
Submitted (26-JUL-1999) CIBC, Royal Ontario Museum, 100 Queen's  
Park, Toronto, Ontario M5S 2C6, Canada  
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DB 82 GTTATTTCGTGAAGTCTGAAGAG 59

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DEFINITION  
ACCESSION  
VERSION  
KEYWORDS  
SOURCE  
ORGANISM  
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AUTHORS  
TITLE  
JOURNAL  
MEDLINE  
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AUTHORS  
TITLE  
JOURNAL  
FEATURES  
source  
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mrna  
BASE COUNT  
ORIGIN

AF174514 100 bp DNA linear VRT 20-APR-2000  
Bufo melanoscticus KIZ-97L118 cytochrome b gene, partial cds;  
mitochondrial gene for mitochondrial product.  
AF174514  
AF174514.1 GI:7620465  
Bufo melanoscticus.  
Mitochondrion Bufo melanoscticus  
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Amphibia; Batrachia; Anura; Neobatrachia; Bufonidae;  
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1 (bases 1 to 100)  
Liu, W., Lathrop, A., Fu, J., Yang, D. and Murphy, R.W.  
Phylogeny of East Asian bufonids inferred from mitochondrial DNA  
sequences (Anura: Amphibia)  
Mol. Phylogenet. Evol. 14 (3), 423-435 (2000)

2 (bases 1 to 100)  
Liu, W., Lathrop, A., Fu, J. and Murphy, R.W.  
Direct Submission  
Submitted (26-JUL-1999) CIBC, Royal Ontario Museum, 100 Queen's  
Park, Toronto, Ontario M5S 2C6, Canada  
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DB 82 GTTATTTCGTGAAGTCTGAAGAG 59

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DEFINITION  
ACCESSION  
VERSION  
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ORGANISM  
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TITLE  
JOURNAL  
MEDLINE  
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AUTHORS  
TITLE  
JOURNAL  
FEATURES  
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CDS  
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BASE COUNT  
ORIGIN

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mitochondrial gene for mitochondrial product.  
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AF174515.1 GI:7620467  
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Mitochondrion Bufo melanoscticus  
Eukaryota; Metazoa; Chordata; Vertebrata; Euteleostomi;  
Amphibia; Batrachia; Anura; Neobatrachia; Bufonidae;  
Bufo.  
1 (bases 1 to 100)  
Liu, W., Lathrop, A., Fu, J., Yang, D. and Murphy, R.W.  
Phylogeny of East Asian bufonids inferred from mitochondrial DNA  
sequences (Anura: Amphibia)  
Mol. Phylogenet. Evol. 14 (3), 423-435 (2000)

2 (bases 1 to 100)  
Liu, W., Lathrop, A., Fu, J. and Murphy, R.W.  
Direct Submission  
Submitted (26-JUL-1999) CIBC, Royal Ontario Museum, 100 Queen's  
Park, Toronto, Ontario M5S 2C6, Canada  
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Matches 10; Conservative 5; Mismatches 9; Indels 0; Gaps 0;

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| : : : : | : | | | | | | | | |  
Db 82 GTTATTTCGTGTAGCCTTAAGAG 59

RESULT 7  
AF174516/c 100 bp DNA linear VRT 20-APR-2000  
LOCUS Bufo melanostictus K12-97L372 cytochrome b gene, partial cds;  
DEFINITION mitochondrial gene for mitochondrial product.  
ACCESSION AF174516  
VERSION AF174516.1 GI:7620469  
KEYWORDS  
SOURCE Bufo melanostictus.  
ORGANISM Mitochondrion Bufo melanostictus  
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
Amphibia; Batrachia; Anura; Neobatrachia; Bufonidae; Bufonidae;  
Bufo.

REFERENCE 1 (bases 1 to 100)  
AUTHORS Liu, W., Lathrop, A., Fu, J., Yang, D. and Murphy, R. W.  
TITLE Phylogeny of East Asian bufonids inferred from mitochondrial DNA  
sequences (Anura: Amphibia)  
JOURNAL Mol. Phylogenet. Evol. 14 (3), 423-435 (2000)  
MEDLINE 20179527  
PUBMED 10712847

REFERENCE 2 (bases 1 to 100)  
AUTHORS Liu, W., Lathrop, A., Fu, J. and Murphy, R. W.  
TITLE Direct Submission  
JOURNAL Submitted (26-JUL-1999) CBCB, Royal Ontario Museum, 100 Queen's  
Park, Toronto, Ontario M5S 2C6, Canada  
Location/Qualifiers

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Db 82 GTTATTTCGTGTAGCCTTAAGAG 59

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DEFINITION mitochondrial gene for mitochondrial product.  
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VERSION AF174517.1 GI:7620471  
KEYWORDS  
SOURCE Bufo melanostictus.  
ORGANISM Mitochondrion Bufo melanostictus

REFERENCE 1 (bases 1 to 100)  
AUTHORS Liu, W., Lathrop, A., Fu, J., Yang, D. and Murphy, R. W.  
TITLE Phylogeny of East Asian bufonids inferred from mitochondrial DNA  
sequences (Anura: Amphibia)  
JOURNAL Mol. Phylogenet. Evol. 14 (3), 423-435 (2000)  
MEDLINE 20179527  
PUBMED 10712847

REFERENCE 2 (bases 1 to 100)  
AUTHORS Liu, W., Lathrop, A., Fu, J. and Murphy, R. W.  
TITLE Direct Submission  
JOURNAL Submitted (26-JUL-1999) CBCB, Royal Ontario Museum, 100 Queen's  
Park, Toronto, Ontario M5S 2C6, Canada  
Location/Qualifiers

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DEFINITION mitochondrial gene for mitochondrial product.  
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VERSION AF174518.1 GI:7620473  
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SOURCE Bufo melanostictus.  
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Amphibia; Batrachia; Anura; Neobatrachia; Bufonidae; Bufonidae;  
Bufo.

REFERENCE 1 (bases 1 to 100)  
AUTHORS Liu, W., Lathrop, A., Fu, J., Yang, D. and Murphy, R. W.  
TITLE Phylogeny of East Asian bufonids inferred from mitochondrial DNA  
sequences (Anura: Amphibia)  
JOURNAL Mol. Phylogenet. Evol. 14 (3), 423-435 (2000)  
MEDLINE 20179527  
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REFERENCE 2 (bases 1 to 100)  
AUTHORS Liu, W., Lathrop, A., Fu, J. and Murphy, R. W.  
TITLE Direct Submission  
JOURNAL Submitted (26-JUL-1999) CBCB, Royal Ontario Museum, 100 Queen's  
Park, Toronto, Ontario M5S 2C6, Canada  
Location/Qualifiers

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82 GTTGTCTTCTGTGAGCCCTAAGAG 59

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LOCUS  
DEFINITION Bufo melanostictus ROM 33163 cytochrome b gene, partial cds;  
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ACCESSION  
VERSION AF174519.1 GI:7620475  
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ORGANISM Bufo melanostictus.  
Mitochondrion Bufo melanostictus  
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Amphibia; Batrachia; Anura; Neobatrachia; Bufonidae; Bufonidae;  
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1 (bases 1 to 100)  
Liu, W., Lathrop, A., Fu, J., Yang, D. and Murphy, R.W.  
Phylogeny of East Asian bufonids inferred from mitochondrial DNA  
sequences (Anura: Amphibia)  
Mol. Phylogenet. Evol. 14 (3), 423-435 (2000)  
20179527  
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Liu, W., Lathrop, A., Fu, J. and Murphy, R.W.  
Direct Submission  
Submitted (26-JUL-1999) CIBCB, Royal Ontario Museum, 100 Queen's  
Park, Toronto, Ontario M5S 2C6, Canada  
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Db 82 GTTGTCTTCTGTGAGCCCTAAGAG 59

RESULT 11 100 bp DNA linear VRT 20-APR-2000  
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LOCUS  
DEFINITION Bufo melanostictus ROM 33855 cytochrome b gene, partial cds;  
AF174520  
ACCESSION  
VERSION AF174520.1 GI:7620477  
KEYWORDS  
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ORGANISM Bufo melanostictus.  
Mitochondrion Bufo melanostictus  
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
Amphibia; Batrachia; Anura; Neobatrachia; Bufonidae; Bufonidae;  
Bufo.  
1 (bases 1 to 100)  
Liu, W., Lathrop, A., Fu, J., Yang, D. and Murphy, R.W.  
Phylogeny of East Asian bufonids inferred from mitochondrial DNA  
sequences (Anura: Amphibia)  
Mol. Phylogenet. Evol. 14 (3), 423-435 (2000)  
20179527  
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2 (bases 1 to 100)  
Liu, W., Lathrop, A., Fu, J. and Murphy, R.W.  
Direct Submission  
Submitted (26-JUL-1999) CIBCB, Royal Ontario Museum, 100 Queen's  
Park, Toronto, Ontario M5S 2C6, Canada  
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Matches 10; Conservative 5; Mismatches 9; Indels 0; Gaps 0;

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DEFINITION Bufo melanostictus ROM 33861 cytochrome b gene, partial cds;  
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VERSION AF174521.1 GI:7620479  
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Mitochondrion Bufo melanostictus  
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Amphibia; Batrachia; Anura; Neobatrachia; Bufonidae; Bufonidae;  
Bufo.

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REFERENCE AUTHORS Liu,W., Lathrop,A., Fu,J., Yang,D. and Murphy,R.W.  
TITLE Phylogeny of East Asian bufonids inferred from mitochondrial DNA  
JOURNAL sequences (Anura: Amphibia)  
MEDLINE MoI. Phylogenet. Evol. 14 (3), 423-435 (2000)  
PUBMED 10719527  
DOI 10.1071/2847  
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REFERENCE AUTHORS Liu,W., Lathrop,A., Fu,J. and Murphy,R.W.  
TITLE Direct Submission  
JOURNAL Submitted (26-Jul-1999) CCBG, Royal Ontario Museum, 100 Queen's  
Park, Toronto, Ontario M5S 2C6, Canada  
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ACCESSION AFI144669  
VERSION AFI144669.1 GI:5690267  
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ORGANISM Patella vulgata  
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Patelloidea; Patelidae; Patella.  
1 (bases 1 to 82)  
de Rosa,R., Grenier,J.R., Andreeva,T., Cook,C.E., Adoutte,A.,  
Akam,M., Carroll,S.B. and Balavoine,G.  
Hox genes in brachiopods and priapulids and protosome evolution  
Nature 399 (6738), 772-776 (1999)  
MEDLINE 99318125  
PUBMED 10391241  
2 (bases 1 to 82)  
de Rosa,R., Larilliot,N. and Adoutte,A.  
Direct Submission  
Submitted (21-Apr-1999) Centre de Genetique Moleculaire, Avenue de  
la terrasse, Gif-sur-Yvette 91198, France  
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DEFINITION	M.domesticus DBA/2 rearranged T-cell receptor (Vgamma2-N-Jgamma2).				
ACCESSION	X63580				
VERSION	X63580.1	GI:57892			
KEYWORDS	Joining region; N-region; T-cell receptor; variable region.				
SOURCE	western European house mouse.				
ORGANISM	Mus musculus domesticus				
REFERENCE	Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.				
AUTHORS	1 (bases 1 to 53)				
JOURNAL	Roger,T.R.				
REFERENCE	Unpublished				
AUTHORS	2 (bases 1 to 53)				
TITLE	Roger,T.				
JOURNAL	Direct Submission				
FEATURES	Submitted (16-DEC-1991) T. Roger, Laboratoire d'immunodifférenciation, Service du Pr SEMAN, Institut J.MONOD, 2, Place JUSSTIER, 75251 PARIS CEDEX 05, FRANCE				
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DEFINITION	M.domesticus DBA/2 rearranged T-cell receptor (Vgamma2-N-Jgamma2).				
ACCESSION	X63580				
VERSION	X63580.1	GI:57892			
KEYWORDS	Joining region; N-region; T-cell receptor; variable region.				
SOURCE	western European house mouse.				
ORGANISM	Mus musculus domesticus				
REFERENCE	Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.				
AUTHORS	1 (bases 1 to 53)				
JOURNAL	Roger,T.R.				
REFERENCE	Unpublished				
AUTHORS	2 (bases 1 to 53)				
TITLE	Roger,T.				
JOURNAL	Direct Submission				
FEATURES	Submitted (16-DEC-1991) T. Roger, Laboratoire d'immunodifférenciation, Service du Pr SEMAN, Institut J.MONOD, 2, Place JUSSTIER, 75251 PARIS CEDEX 05, FRANCE				
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Best Local Similarity	52.4%; Pred. No. 5.3e+03;				
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LOCUS	MDTRVNB/C	53 bp	mRNA	linear	ROD 07-MAR-1993
DEFINITION	M.domesticus DBA/2 rearranged T-cell receptor (Vgamma2-N-Jgamma2).				
ACCESSION	X63580				
VERSION	X63580.1	GI:57892			
KEYWORDS	Joining region; N-region; T-cell receptor; variable region.				
SOURCE	western European house mouse.				
ORGANISM	Mus musculus domesticus				
REFERENCE	Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.				
AUTHORS	1 (bases 1 to 53)				
JOURNAL	Roger,T.R.				
REFERENCE	Unpublished				
AUTHORS	2 (bases 1 to 53)				
TITLE	Roger,T.				
JOURNAL	Direct Submission				
FEATURES	Submitted (16-DEC-1991) T. Roger, Laboratoire d'immunodifférenciation, Service du Pr SEMAN, Institut J.MONOD, 2, Place JUSSTIER, 75251 PARIS CEDEX 05, FRANCE				
LOCUS	Location/Qualifiers				
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LOCUS	MDTRVNB/C	53 bp	mRNA	linear	ROD 07-MAR-1993
DEFINITION	M.domesticus DBA/2 rearranged T-cell receptor (Vgamma2-N-Jgamma2).				
ACCESSION	X63580				
VERSION	X63580.1	GI:57892			
KEYWORDS	Joining region; N-region; T-cell receptor; variable region.				
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REFERENCE	Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.				
AUTHORS	1 (bases 1 to 53)				
JOURNAL	Roger,T.R.				
REFERENCE	Unpublished				
AUTHORS	2 (bases 1 to 53)				
TITLE	Roger,T.				
JOURNAL	Direct Submission				
FEATURES	Submitted (16-DEC-1991) T. Roger, Laboratoire d'immunodifférenciation, Service du Pr SEMAN, Institut J.MONOD, 2, Place JUSSTIER, 75251				



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DEFINITION Bufo himalayanus K12-95L010 cytochrome b gene, partial cds;
ACCESSION AF174501
VERSION AF174501.1 GI:7620439
KEYWORDS
SOURCE Bufo himalayanus.
ORGANISM Mitochondrion Bufo himalayanus
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Amphibia; Batrachia; Anura; Neobatrachia; Bufonidae; Bufonidae;
Bufo.
REFERENCE 1 (bases 1 to 100)
AUTHORS Liu, W., Lathrop, A., Fu, J., Yang, D. and Murphy, R.W.
TITLE Phylogeny of East Asian bufonids inferred from mitochondrial DNA
SEQUENCES (Anura: Amphibia)
JOURNAL Mol. Phylogenet. Evol. 14 (3), 423-435 (2000)
MEDLINE 20179527
PUBMED 10712847
REFERENCE 2 (bases 1 to 100)
AUTHORS Liu, W., Lathrop, A., Fu, J. and Murphy, R.W.
TITLE Direct Submission
JOURNAL Submitted (26-JUL-1999) CBCB, Royal Ontario Museum, 100 Queen's
Park, Toronto, Ontario M5S 2C6, Canada
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Best Local Similarity 40.9%; Pred. No. 5.6e+03;
Matches 9; Conservative 5; Mismatches 8; Indels 0; Gaps 0;
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GenCore version 4.5  
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OM nucleic - nucleic search, using sw model

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Sequence: 1 nnnnauuncununguagcccnangnngn 29

Scoring table: IDENTITY NUC  
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Searched: 1736436 seqs, 858457221 residues 2046006

Total number of hits satisfying chosen parameters:  
Minimum DB seq length: 0  
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Post-processing: Minimum Match 0%  
Maximum Match 100%  
Listing first 45 summaries

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Pred. No. is the number of results predicted by chance to have a  
score greater than or equal to the score of the result being printed,  
and is derived by analysis of the total score distribution.

#### SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	18	62.1	29	21	AAA70827
2	18	62.1	29	21	AAA70828
3	18	62.1	29	21	AAA70829
4	18	62.1	29	21	AAA70830
5	18	62.1	42	21	AAA71113
6	18	62.1	42	21	AAA71114
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18	18	62.1	42	21	AAA71129	Molecular Interact
19	18	62.1	42	21	AAA71131	Molecular Interact
20	18	62.1	42	21	AAA71132	Molecular Interact
21	18	62.1	44	21	AAA71112	Molecular Interact
22	18	62.1	44	21	AAA71125	Molecular Interact
23	18	62.1	44	21	AAA71133	Molecular Interact
24	18	62.1	45	21	AAA70824	Molecular Interact
25	18	62.1	45	21	AAA70825	Molecular Interact
26	18	62.1	45	21	AAA70826	Molecular Interact
27	18	62.1	46	21	AAA71085	Molecular Interact
28	18	62.1	46	21	AAA71087	Molecular Interact
29	18	62.1	46	21	AAA71088	Molecular Interact
30	18	62.1	46	21	AAA71089	Molecular Interact
31	18	62.1	46	21	AAA71090	Molecular Interact
32	18	62.1	46	21	AAA71093	Molecular Interact
33	18	62.1	46	21	AAA71094	Molecular Interact
34	18	62.1	46	21	AAA71095	Molecular Interact
35	18	62.1	46	21	AAA71096	Molecular Interact
36	18	62.1	46	21	AAA71099	Molecular Interact
37	18	62.1	46	21	AAA71100	Molecular Interact
38	18	62.1	46	21	AAA71103	Molecular Interact
39	18	62.1	46	21	AAA71104	Molecular Interact
40	18	62.1	46	21	AAA71105	Molecular Interact
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#### ALIGNMENTS

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AC AAA70827;  
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DF 27-APR-2001 (first entry)  
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DE Molecular Interaction site RNA #27.  
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KW Modulator; Identification; molecular interaction; virtual library; ss.  
XX  
OS Synthetic.  
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PN WO958947-A2.  
XX  
PD 18-NOV-1999.  
XX  
PR 12-MAY-1999; 99WO-US10361.  
XX  
PR 12-MAY-1998; 98US-0076404.  
XX  
PR 12-MAY-1998; 98US-0085092.  
XX  
PA (ISIS-) ISIS PHARM, INC.  
XX  
PI Ecker DJ, Grifey R, Crooke ST, Sampath R, Swayze E, Mohan V;  
XX  
PI Hostadler S, McNeil J;  
XX  
DR WPI: 2000-086439/07.  
XX  
PT Identifying compounds which modulate activity of target biomolecules,  
PT used to provide compounds which can be used as pharmacological,

PT agricultural and industrial compounds -  
XX  
PS Claim 235; Page 235; 405pp; English.  
XX  
CC This invention describes a novel method for identifying compounds which  
CC modulate the activity of a target biomolecule. The method uses  
CC 3-dimensional representations of the biomolecule and a library of  
CC compounds and comprises (a) identifying at least one molecular  
CC interaction site of the target RNA; (b) generating in silico a virtual  
CC library of compounds predicted or calculated to interact with the  
CC molecular interaction site; and (c) comparing 3-dimensional (3-D)  
CC representations of the target RNA with members of the virtual library of  
CC compounds to generate a hierarchy of the compounds ranked in accordance  
CC with their respective ability to form physical interactions with the  
CC molecular interaction site. The method also describes (1) RNA comprising  
CC a joined sequence of at least 24 nucleotides but not more than 70  
CC nucleotides and having secondary structure defined by: (a) 3 nucleotides  
CC forming a first side of a first double stranded (ds) region; (b) 2  
CC nucleotides forming a first side of an internal loop region; (c) 4  
CC nucleotides forming a first side of a second ds region; (d) 4 or 5  
CC nucleotides forming an end loop region; (e) 4 nucleotides forming a  
CC second side of the second ds region; (f) 4 nucleotides forming a second  
CC side of the internal loop region; and (g) 3 nucleotides forming a second  
CC side of the first ds region; (2) a purified and isolated RNA fragment  
CC comprising the human sequence UUUACACUAUUCUGUUACGAAAUUC (11). The  
CC methods and products can be used for identifying agents which modulate  
CC the activity of biomolecules, particularly RNA. Such agents can be used  
CC as pharmaceutical, agricultural or industrial compounds.  
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PN WO958947-A2.  
XX  
PD 18-NOV-1999.  
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PF 12-MAY-1999; 99WO-US10361.  
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PR 12-MAY-1998; 98US-0076404.  
XX  
PR 12-MAY-1998; 98US-0085092.  
XX  
PA (ISIS-) ISIS PHARM INC.  
XX  
PI Ecker DJ, Griffey R, Crooke ST, Sampath R, Swayze E, Mohan V;  
XX  
PI Hofstadler S, McNeil J;  
XX  
DR WPI; 2000-086439/07.  
XX  
PT Identifying compounds which modulate activity of target biomolecules,  
PT used to provide compounds which can be used as pharmacological,

PT agricultural and industrial compounds -  
XX  
PS Claim 235; Page 235; 405pp; English.  
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CC This invention describes a novel method for identifying compounds which  
CC modulate the activity of a target biomolecule. The method uses  
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CC compounds and comprises (a) identifying at least one molecular  
CC interaction site of the target RNA; (b) generating in silico a virtual  
CC library of compounds predicted or calculated to interact with the  
CC molecular interaction site; and (c) comparing 3-dimensional (3-D)  
CC representations of the target RNA with members of the virtual library of  
CC compounds to generate a hierarchy of the compounds ranked in accordance  
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CC a joined sequence of at least 24 nucleotides but not more than 70  
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CC nucleotides forming a first side of an internal loop region; (c) 4  
CC nucleotides forming a first side of a second ds region; (d) 4 or 5  
CC nucleotides forming an end loop region; (e) 4 nucleotides forming a  
CC second side of the second ds region; (f) 4 nucleotides forming a second  
CC side of the internal loop region; and (g) 3 nucleotides forming a second  
CC side of the first ds region; (2) a purified and isolated RNA fragment  
CC comprising the human sequence UUUACACUAUUCUGUUACGAAAUUC (11). The  
CC methods and products can be used for identifying agents which modulate  
CC the activity of biomolecules, particularly RNA. Such agents can be used  
CC as pharmaceutical, agricultural or industrial compounds.  
XX  
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Best Local Similarity 75.0%; Pred. No. 0.97;  
Matches 18; Conservative 0; Mismatches 6; Indels 0; Gaps 0;  
  
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Db 4 gauncuuuunguaagcccnaggg 27  
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OS Mus sp.  
XX  
PN WO958947-A2.  
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PD 18-NOV-1999.  
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PF 12-MAY-1999; 99WO-US10361.  
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PR 12-MAY-1998; 98US-0076404.  
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PR 12-MAY-1998; 98US-0085092.  
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PI Hofstadler S, McNeil J;  
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DR WPI; 2000-086439/07.  
XX  
PT Identifying compounds which modulate activity of target biomolecules,  
PT used to provide compounds which can be used as pharmacological,

PT agricultural and industrial compounds -  
XX  
PS Claim 235; Page 235; 405pp; English.  
XX  
CC This invention describes a novel method for identifying compounds which  
CC modulate the activity of a target biomolecule. The method uses  
CC 3-dimensional representations of the biomolecule and a library of  
CC compounds and comprises (a) identifying at least one molecular  
CC interaction site of the target RNA; (b) generating in silico a virtual  
CC library of compounds predicted or calculated to interact with the  
CC molecular interaction site; and (c) comparing 3-dimensional (3-D)  
CC representations of the target RNA with members of the virtual library of  
CC compounds to generate a hierarchy of the compounds ranked in accordance  
CC with their respective ability to form physical interactions with the  
CC molecular interaction site. The method also describes (1) RNA comprising  
CC a joined sequence of at least 24 nucleotides but not more than 70  
CC nucleotides and having secondary structure defined by: (a) 3 nucleotides  
CC forming a first side of a first double stranded (ds) region; (b) 2  
CC nucleotides forming a first side of an internal loop region; (c) 4  
CC nucleotides forming a first side of a second ds region; (d) 4 or 5  
CC nucleotides forming an end loop region; (e) 4 nucleotides forming a  
CC second side of the second ds region; (f) 4 nucleotides forming a  
CC side of the internal loop region; and (g) 3 nucleotides forming a second  
CC side of the first ds region; (2) a purified and isolated RNA fragment  
CC comprising the human sequence UUUACACAUUAUCUAGUUCACGAAAUUC (II). The  
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CC the activity of biomolecules, particularly RNA. Such agents can be used  
CC as pharmaceutical, agricultural or industrial compounds.  
SQ Sequence 29 BP; 8 A; 6 C; 6 G; 9 U; 0 other;

Query Match 62.1%; Score 18; DB 21; Length 29;  
Best Local Similarity 75.0%; Pred. No. 0.97;  
Matches 18; Conservative 0; Mismatches 6; Indels 0; Gaps 0;  
Qy 4 gauncuununguaagcccnangng 27  
||| |||| |||| |||| |||  
Db 4 gauncuununguaagcccaagsg 27

RESULT 4  
ID AAA70830 standard; RNA; 29 BP.  
XX  
AC AAA70830;  
XX  
DT 27-APR-2001 (first entry)  
XX  
XX Molecular interaction site RNA #30.  
KM Modulator; identification; molecular interaction; virtual library; ss.  
XX  
OS Rattus sp.  
XX  
PN WO9558947-A2.  
XX  
PD 18-NOV-1999.  
XX  
PF 12-MAY-1999; 99WO-US10361.  
XX  
PR 12-MAY-1998; 98US-0076404.  
PR 12-MAY-1998; 98US-0085092.  
XX  
PA (ISIS-) ISIS PHARM INC.  
XX  
PI Ecker DJ, Griffey R, Crooke ST, Sampath R, Swayze E, Mohan V;  
PI Hofstadler S, McNeil J;  
XX  
DR WPI; 2000-086439/07.  
XX  
PT Identifying compounds which modulate activity of target biomolecules,  
PT used to provide compounds which can be used as pharmacological,

PT agricultural and industrial compounds -  
XX  
PS Claim 235; Page 235; 405pp; English.  
XX  
CC This invention describes a novel method for identifying compounds which  
CC modulate the activity of a target biomolecule. The method uses  
CC 3-dimensional representations of the biomolecule and a library of  
CC compounds and comprises (a) identifying at least one molecular  
CC interaction site of the target RNA; (b) generating in silico a virtual  
CC library of compounds predicted or calculated to interact with the  
CC molecular interaction site; and (c) comparing 3-dimensional (3-D)  
CC representations of the target RNA with members of the virtual library of  
CC compounds to generate a hierarchy of the compounds ranked in accordance  
CC with their respective ability to form physical interactions with the  
CC molecular interaction site. The method also describes (1) RNA comprising  
CC a joined sequence of at least 24 nucleotides but not more than 70  
CC nucleotides and having secondary structure defined by: (a) 3 nucleotides  
CC forming a first side of a first double stranded (ds) region; (b) 2  
CC nucleotides forming a first side of an internal loop region; (c) 4  
CC nucleotides forming a first side of a second ds region; (d) 4 or 5  
CC nucleotides forming an end loop region; (e) 4 nucleotides forming a  
CC second side of the second ds region; (f) 4 nucleotides forming a  
CC side of the internal loop region; and (g) 3 nucleotides forming a second  
CC side of the first ds region; (2) a purified and isolated RNA fragment  
CC comprising the human sequence UUUACACAUUAUCUAGUUCACGAAAUUC (II). The  
CC methods and products can be used for identifying agents which modulate  
CC the activity of biomolecules, particularly RNA. Such agents can be used  
CC as pharmaceutical, agricultural or industrial compounds.  
SQ Sequence 29 BP; 8 A; 6 C; 6 G; 9 U; 0 other;

Query Match 62.1%; Score 18; DB 21; Length 29;  
Best Local Similarity 75.0%; Pred. No. 0.97;  
Matches 18; Conservative 0; Mismatches 6; Indels 0; Gaps 0;  
Qy 4 gauncuununguaagcccnangng 27  
||| |||| |||| |||| |||  
Db 4 gauncuununguaagcccaagsg 27

RESULT 5  
ID AAA71113 standard; RNA; 42 BP.  
XX  
AC AAA71113;  
XX  
DT 27-APR-2001 (first entry)  
XX  
XX Molecular interaction site RNA #189.  
DE Modulator; identification; molecular interaction; virtual library; ss.  
XX  
XX unidentified.  
XX  
PN WO9558947-A2.  
XX  
PD 18-NOV-1999.  
XX  
PF 12-MAY-1999; 99WO-US10361.  
XX  
PR 12-MAY-1998; 98US-0076404.  
PR 12-MAY-1998; 98US-0085092.  
XX  
PA (ISIS-) ISIS PHARM INC.  
XX  
PI Ecker DJ, Griffey R, Crooke ST, Sampath R, Swayze E, Mohan V;  
PI Hofstadler S, McNeil J;  
XX  
DR WPI; 2000-086439/07.  
XX  
PT Identifying compounds which modulate activity of target biomolecules,  
PT used to provide compounds which can be used as pharmacological,

PT - agricultural and industrial compounds  
XX  
PS Example 7; Figure 122; 405pp; English.

This invention describes a novel method for identifying compounds which modulate the activity of a target biomolecule. The method uses 3-dimensional representations of the biomolecule and a library of compounds and comprises (a) identifying at least one molecular interaction site of the target RNA; (b) generating in silico a virtual library of compounds predicted or calculated to interact with the molecular interaction site; and (c) comparing 3-dimensional (3-D) representations of the target RNA with members of the virtual library of compounds to generate a hierarchy of the compounds ranked in accordance with their respective ability to form physical interactions with the molecular interaction site. The method also describes (1) RNA comprising a joined sequence of at least 24 nucleotides but not more than 70 nucleotides and having secondary structure defined by: (a) 3 nucleotides forming a first side of a first double stranded (ds) region; (b) 2 nucleotides forming a first side of an internal loop region; (c) 4 nucleotides forming a first side of a second ds region; (d) 4 or 5 nucleotides forming an end loop region; (e) 4 nucleotides forming a second side of the second ds region; (f) 4 nucleotides forming a second side of the internal loop region; and (g) 3 nucleotides forming a second side of the first ds region; (2) a purified and isolated RNA fragment comprising the human sequence GUUACACUAAUUGAGUUUACGAAAAAC (II). The methods and products can be used for identifying agents which modulate the activity of biomolecules, particularly RNA. Such agents can be used as pharmaceutical, agricultural or industrial compounds.

Query Match	62.1%	Score 18	DB 21	Length 42
Best Local Similarity	75.0%	Pred. No. 1		
Matches 18	Conservative 0	Mismatches 6	Indels 0	Gaps 0

Qy 4 gauncuuunguaagcccnangng 27  
 ||| ||| ||| ||| | | |  
 Db 7 gaucuuuunguaagcccuacng 30

RESULT	6
AAA71114	
ID	AAA71114 standard; RNA; 42 BP

27-APR-2001 (first entry)

DE Molecular interaction site RNA #190.

KW modulator; identification; molecular interaction; virtual library; ss

OS Unidentified

PN W09958947-A2

PD 18-NOV-1999.

PF 12-MAY-1999; 99WO-US10361

PR 12-MAY-1998; 98US-0076404

PR 12-MAY-1998; 98US-0085092

PA (ISIS-) ISIS PHARM INC.

PI Ecker DJ, Griffey R, Crooke ST, Sampath R, Swayze E, Mohan V.

PI Hofstadler S, McNeill J;

DR WPI; 2000-086439/07.

PT Identifying compounds which modulate activity of target biomolecules  
PT used to provide compounds which can be used as pharmacological,

PT	agricultural and industrial compounds
XX	
PS	Example 7; Figure 122; 405pp; English.

This invention describes a novel method for identifying compounds which modulate the activity of a target biomolecule. The method uses 3-dimensional representations of the biomolecule and a library of compounds and comprises (a) identifying at least one molecular interaction site of the target RNA; (b) generating in silico a virtual library of compounds predicted or calculated to interact with the molecular interaction site; and (c) comparing 3-dimensional (3-D) representations of the target RNA with members of the virtual library of compounds to generate a hierarchy of the compounds ranked in accordance with their respective ability to form physical interactions with the molecular interaction site. The method also describes (1) RNA comprising a joined sequence of at least 24 nucleotides but not more than 70 nucleotides and having secondary structure defined by: (a) 3 nucleotides forming a first side of a first double stranded (ds) region; (b) 2 nucleotides forming a first side of an internal loop region; (c) 4 nucleotides forming a first side of a second ds region; (d) 4 or 5 nucleotides forming an end loop region; (e) 4 nucleotides forming a second side of the second ds region; (f) 4 nucleotides forming a second side of the internal loop region; and (g) 3 nucleotides forming a second side of the first ds region; (2) a purified and isolated RNA fragment comprising the human sequence UUUACAGUAAUUNUGUUGUACGAAAAAC (II). The methods and products can be used for identifying agents which modulate the activity of biomolecules, particularly RNA. Such agents can be used as pharmaceutical, agricultural or industrial compounds.

Query Match	62.1%	Score 18;	DB 21;	Length 42;
Best Local Similarity	75.0%	Pred. No. 1;		
Matches 18; Conservative	0;	Mismatches 6;	Indels 0;	Gaps 0;

**DY**      4    gauncuuunguaagcccnang    27  
           ||| |||| | ||||| | |  
**Db**      7    gaucuuuuuguaagcccuagcg    30

RESULT	7
AAA71115	
ID	AAA71115 standard; RNA; 42 BP

DT 27-APR-2001 (first entry)

DE molecular interaction site RNA #191.

KW modulator; identification; molecular interaction; virtual library; ss

OS Unidentified

PN W09958947-A2

PD 18-NOV-1999

PF 12-MAY-1999; 99WO-US10361

PR 12-MAY-1998; 98US-0076404

PR 12-MAY-1998; 98US-0085092

PA (ISIS-) ISIS PHARM INC.

PI Ecker DJ, Griffey R, Crooke ST, Sampath R, Swayze E, Mohan V,

PI Hofstadler S, McNeil J;

DR WPI; 2000-086439/07.

PT Identifying compounds which modulate activity of target biomolecules  
PT used to provide compounds which can be used as pharmacological,

PT Identifying compounds which modulate activity of target biomolecules  
PT used to provide compounds which can be used as pharmacological,

PT agricultural and industrial compounds -  
XX  
XX  
XX Example 7; Figure 125; 405bp; English.  
CC  
CC This invention describes a novel method for identifying compounds which  
CC modulate the activity of a target biomolecule. The method uses  
CC 3-dimensional representations of the biomolecule and a library of  
CC compounds and comprises (a) identifying at least one molecular  
CC interaction site of the target RNA; (b) generating in silico a virtual  
CC library of compounds predicted or calculated to interact with the  
CC molecular interaction site; and (c) comparing 3-dimensional (3-D)  
CC representations of the target RNA with members of the virtual library of  
CC compounds to generate a hierarchy of the compounds ranked in accordance  
CC with their respective ability to form physical interactions with the  
CC molecular interaction site. The method also describes (1) RNA comprising  
CC a joined sequence of at least 24 nucleotides but not more than 70  
CC nucleotides and having secondary structure defined by: (a) 3 nucleotides  
CC forming a first side of a first double stranded (ds) region; (b) 2  
CC nucleotides forming a first side of an internal loop region; (c) 4  
CC nucleotides forming a first side of a second ds region; (d) 4 or 5  
CC nucleotides forming an end loop region; (e) 4 nucleotides forming a  
CC second side of the second ds region; (f) 4 nucleotides forming a second  
CC side of the internal loop region; and (g) 3 nucleotides forming a second  
CC side of the first ds region; (2) a purified and isolated RNA fragment  
CC comprising the human sequence UUUACACUUAUUCUGUUGUACGAAAUUC (II). The  
CC methods and products can be used for identifying agents which modulate  
CC the activity of biomolecules, particularly RNA. Such agents can be used  
CC as pharmaceutical, agricultural or industrial compounds.  
XX  
XX Sequence 42 BP; 12 A; 7 C; 6 G; 17 T; 0 other;  
50

Query Match	62.18;	Score 18;	DB 21;	Length 42;
Best Local Similarity	54.28;	Pred. NO. 1;		
Matches 13; Conservative	5;	Mismatches 6;	Indels 0;	Gaps 0;

4 gauncuunguaagcccnang 27  
 7 gattcttltgtaagccctacgy 30

```

RESULT 10
AAA71119
ID AAA71119 standard; DNA; 42 BP
xx
AC AAA71119;

```

27-APR-2001 (flrst entry)

DE Molecular interaction site DNA #125.

KW Modulator; identification; molecular interaction; virtual library; ss.

OS Unidentified

PN W09958947-A2

PD 18-NOV-1999.

PF 12-MAY-1999;

PR 12-MAY-1998; 98US-0076404.

XX

XX

PI Hofstadler S, McNeill J;

DR WPI; 2000-086439/07.

PT Identifying compounds which modulate activity of target biomolecules used to provide compounds which can be used as pharmacological,

PT agricultural and industrial compounds  
XX  
PS Example 7; Figure 125; 405pp; English.

CC This invention describes a novel method for identifying compounds which  
CC modulate the activity of a target biomolecule. The method uses  
CC 3-dimensional representations of the biomolecule and a library of  
CC compounds and comprises (a) identifying at least one molecular  
CC interaction site of the target RNA; (b) generating *in silico* a virtual  
CC library of compounds predicted or calculated to interact with the  
CC molecular interaction site; and (c) comparing 3-dimensional (3-D)  
CC representations of the target RNA with members of the virtual library of  
CC compounds to generate a hierarchy of the compounds ranked in accordance  
CC with their respective ability to form physical interactions with the  
CC molecular interaction site. The method also describes (1) RNA comprising  
CC a joined sequence of at least 24 nucleotides but not more than 70  
CC nucleotides and having secondary structure defined by: (a) 3 nucleotides  
CC forming a first side of a first double stranded (ds) region; (b) 2  
CC nucleotides forming a first side of an internal loop region; (c) 4  
CC nucleotides forming a first side of a second ds region; (d) 4 or 5  
CC nucleotides forming an end loop region; (e) 4 nucleotides forming a  
CC second side of the second ds region; (f) 4 nucleotides forming a second  
CC side of the internal loop region; and (g) 3 nucleotides forming a second  
CC side of the first ds region; (2) a purified and isolated RNA fragment  
CC comprising the human sequence UUUUACACAAUUCUAGUUUACGAAAUAC (II). The  
CC methods and products can be used for identifying agents which modulate  
CC the activity of biomolecules, particularly RNA. Such agents can be used  
CC as pharmaceutical, agricultural or industrial compounds.

Query Match	62.1%;	Score 18;	DB 21;	Length 42;
Best Local Similarity	54.2%;	Pred. No. 1;		
Matches 13;	Conservative 5;	Mismatches 6;	Indels 0;	Gaps 0;

QY 4 gauncuunnguagcccnangng 27  
||:|::|:||||| | | |  
Db 7 gattcctttgtlaagcccltagcg 30

```
RESULT 11
AAA71120
ID AAA71120 standard; DNA; 42 BP
XX
AC AAA71120;
```

T 27-APR-2001 (first entry)

DE Molecular interaction site DNA #126.

KW Modulator; identification; molecular interaction; virtual library; ss.

OS Unidentified

PN W09958947-A2

PD 18-NOV-1999.

PF 12-MAY-1999;

PR 12-MAY-1998; 98US-0076404.

XX

XX

PI Hofstadler S, McNeil J;

DR WPI; 2000-086439/07

PT Identifying compounds which modulate activity of target biomolecules, used to provide compounds which can be used as pharmacological, PT



PT agricultural and industrial compounds -  
XX  
PS  
XX Example 7; Figure 125; 405pp; English.  
CC This invention describes a novel method for identifying compounds which  
CC modulate the activity of a target biomolecule. The method uses  
CC 3-dimensional representations of the biomolecule and a library of  
CC compounds and comprises (a) identifying at least one molecular  
CC interaction site of the target RNA; (b) generating in silico a virtual  
CC library of compounds predicted or calculated to interact with the  
CC molecular interaction site; and (c) comparing 3-dimensional (3-D)  
CC representations of the target RNA with members of the virtual library of  
CC compounds to generate a hierarchy of the compounds ranked in accordance  
CC with their respective ability to form physical interactions with the  
CC molecular interaction site. The method also describes (1) RNA comprising  
CC a joined sequence of at least 24 nucleotides but not more than 70  
CC nucleotides and having secondary structure defined by: (a) 3 nucleotides  
CC forming a first side of a first double stranded (ds) region; (b) 2  
CC nucleotides forming a first side of an internal loop region; (c) 4  
CC nucleotides forming a first side of a second ds region; (d) 4 or 5  
CC nucleotides forming an end loop region; (e) 4 nucleotides forming a  
CC second side of the second ds region; (f) 4 nucleotides forming a second  
CC side of the internal loop region; and (g) 3 nucleotides forming a second  
CC side of the first ds region; (2) a purified and isolated RNA fragment  
CC comprising the human sequence UUUACACAUAAUUCUUAUACGAAAUUC (II). The  
CC methods and products can be used for identifying agents which modulate  
CC the activity of biomolecules, particularly RNA. Such agents can be used  
CC as pharmaceutical, agricultural or industrial compounds.  
SQ Sequence 42 BP; 13 A; 7 C; 7 G; 15 T; 0 other;

Query Match 62.1%; Score 18; DB 21; Length 42;  
Best Local Similarity 54.2%; Pred. No. 1;  
Matches 13; Conservative 5; Mismatches 6; Indels 0; Gaps 0;  
QY 4 gauncuununguaagcccnangng 27  
||:|::|:||||| | | |  
DB 7 gatctctttgtgaagcccaagg 30

RESULT 12  
AAA71121  
ID AAA71121 standard; DNA; 42 BP.  
XX  
AC AAA71121;  
XX  
DT 27-APR-2001 (first entry)  
XX  
DE Molecular interaction site DNA #127.  
XX  
KM Modulator; Identification; molecular interaction; virtual library; ss.  
XX  
OS Unidentified.  
XX  
PN WO958947-A2.  
XX  
PD 18-NOV-1999.  
XX  
PF 12-MAY-1999; 99WO-US10361.  
XX  
PR 12-MAY-1998; 98US-0076404.  
XX  
PR 12-MAY-1998; 98US-0085092.  
XX  
PA (ISIS-) ISIS PHARM INC.  
XX  
PI Ecker DJ, Griffey R, Crooke SF, Sampath R, Swayze E, Mohan V;  
PI Hofstadler S, McNeill J;  
XX  
DR WPI; 2000-086439/07.  
XX  
PT Identifying compounds which modulate activity of target biomolecules,  
PT used to provide compounds which can be used as pharmacological,

PT agricultural and industrial compounds -  
XX  
PS  
XX Example 7; Figure 125; 405pp; English.  
CC This invention describes a novel method for identifying compounds which  
CC modulate the activity of a target biomolecule. The method uses  
CC 3-dimensional representations of the biomolecule and a library of  
CC compounds and comprises (a) identifying at least one molecular  
CC interaction site of the target RNA; (b) generating in silico a virtual  
CC library of compounds predicted or calculated to interact with the  
CC molecular interaction site; and (c) comparing 3-dimensional (3-D)  
CC representations of the target RNA with members of the virtual library of  
CC compounds to generate a hierarchy of the compounds ranked in accordance  
CC with their respective ability to form physical interactions with the  
CC molecular interaction site. The method also describes (1) RNA comprising  
CC a joined sequence of at least 24 nucleotides but not more than 70  
CC nucleotides and having secondary structure defined by: (a) 3 nucleotides  
CC forming a first side of a first double stranded (ds) region; (b) 2  
CC nucleotides forming a first side of an internal loop region; (c) 4  
CC nucleotides forming a first side of a second ds region; (d) 4 or 5  
CC nucleotides forming an end loop region; (e) 4 nucleotides forming a  
CC second side of the second ds region; (f) 4 nucleotides forming a second  
CC side of the internal loop region; and (g) 3 nucleotides forming a second  
CC side of the first ds region; (2) a purified and isolated RNA fragment  
CC comprising the human sequence UUUACACAUAAUUCUUAUACGAAAUUC (II). The  
CC methods and products can be used for identifying agents which modulate  
CC the activity of biomolecules, particularly RNA. Such agents can be used  
CC as pharmaceutical, agricultural or industrial compounds.  
SQ Sequence 42 BP; 13 A; 7 C; 7 G; 15 T; 0 other;

Query Match 62.1%; Score 18; DB 21; Length 42;  
Best Local Similarity 54.2%; Pred. No. 1;  
Matches 13; Conservative 5; Mismatches 6; Indels 0; Gaps 0;  
QY 4 gauncuununguaagcccnangng 27  
||:|::|:||||| | | |  
DB 7 gatctctttgtgaagcccaagg 30

RESULT 13  
AAA71123  
ID AAA71123 standard; DNA; 42 BP.  
XX  
AC AAA71123;  
XX  
DT 27-APR-2001 (first entry)  
XX  
DE Molecular interaction site DNA #129.  
XX  
KM Modulator; Identification; molecular interaction; virtual library; ss.  
XX  
OS Unidentified.  
XX  
PN WO958947-A2.  
XX  
PD 18-NOV-1999.  
XX  
PF 12-MAY-1999; 99WO-US10361.  
XX  
PR 12-MAY-1998; 98US-0076404.  
XX  
PR 12-MAY-1998; 98US-0085092.  
XX  
PA (ISIS-) ISIS PHARM INC.  
XX  
PI Ecker DJ, Griffey R, Crooke SF, Sampath R, Swayze E, Mohan V;  
PI Hofstadler S, McNeill J;  
XX  
DR WPI; 2000-086439/07.  
XX  
PT Identifying compounds which modulate activity of target biomolecules,  
PT used to provide compounds which can be used as pharmacological,

PT agricultural and industrial compounds -  
XX  
PS Example 7; Figure 125; 405pp; English.  
XX  
CC This invention describes a novel method for identifying compounds which  
CC modulate the activity of a target biomolecule. The method uses  
CC 3-dimensional representations of the biomolecule and a library of  
CC compounds and comprises (a) identifying at least one molecular  
CC interaction site of the target RNA; (b) generating in silico a virtual  
CC library of compounds predicted or calculated to interact with the  
CC molecular interaction site; and (c) comparing 3-dimensional (3-D)  
CC representations of the target RNA with members of the virtual library of  
CC compounds to generate a hierarchy of the compounds ranked in accordance  
CC with their respective ability to form physical interactions with the  
CC molecular interaction site. The method also describes (1) RNA comprising  
CC a joined sequence of at least 24 nucleotides but not more than 70  
CC nucleotides and having secondary structure defined by: (a) 3 nucleotides  
CC forming a first side of a first double stranded (ds) region; (b) 2  
CC nucleotides forming a first side of an internal loop region; (c) 4  
CC nucleotides forming a first side of a second ds region; (d) 4 or 5  
CC nucleotides forming an end loop region; (e) 4 nucleotides forming a  
CC second side of the second ds region; (f) 4 nucleotides forming a second  
CC side of the internal loop region; and (g) 3 nucleotides forming a second  
CC side of the first ds region; (2) a purified and isolated RNA fragment  
CC comprising the human sequence UUUACACAUACUUGUUCAGAAAUUC (11). The  
CC methods and products can be used for identifying agents which modulate  
CC the activity of biomolecules, particularly RNA. Such agents can be used  
CC as pharmaceutical, agricultural or industrial compounds.  
XX  
SQ Sequence 42 BP; 9 A; 6 C; 9 G; 18 T; 0 other;  
  
Query Match 62.18; Score 18; DB 21; Length 42;  
Best Local Similarity 54.2%; Pred. No. 1;  
Matches 13; Conservative 5; Mismatches 6; Indels 0; Gaps 0;  
  
QY 4 gauncuununguagcccnangng 27  
||:|:::|:||||| | | |  
Db 7 gatcttcttgtaagcctcaggg 30  
  
RESULT 14  
AAA71124  
ID AAA71124 standard; DNA: 42 BP.  
XX  
AC AAA71124;  
  
27-APR-2001 (first entry)  
  
DE Molecular interaction site DNA #130.  
XX  
KM Modulator; identification; molecular interaction; virtual library; ss.  
XX  
OS Unidentified.  
XX  
PN W09958947-A2.  
XX  
PD 18-NOV-1999.  
XX  
PE 12-MAY-1999; 99WO-US10361.  
XX  
PR 12-MAY-1998; 98US-0076404.  
XX  
PR 12-MAY-1998; 98US-0085092.  
XX  
PA (ISIS-) ISIS PHARM INC.  
XX  
PI Ecker DJ, Griffey R, Crooke ST, Sampath R, Swayze E, Mohan V;  
PI Hofstadler S, McNeil J;  
XX  
DR WPI; 2000-086439/07.  
XX  
PT Identifying compounds which modulate activity of target biomolecules,  
PT used to provide compounds which can be used as pharmacological.

PT agricultural and industrial compounds -  
XX  
PS Example 7; Figure 125; 405pp; English.  
XX  
CC This invention describes a novel method for identifying compounds which  
CC modulate the activity of a target biomolecule. The method uses  
CC 3-dimensional representations of the biomolecule and a library of  
CC compounds and comprises (a) identifying at least one molecular  
CC interaction site of the target RNA; (b) generating in silico a virtual  
CC library of compounds predicted or calculated to interact with the  
CC molecular interaction site; and (c) comparing 3-dimensional (3-D)  
CC representations of the target RNA with members of the virtual library of  
CC compounds to generate a hierarchy of the compounds ranked in accordance  
CC with their respective ability to form physical interactions with the  
CC molecular interaction site. The method also describes (1) RNA comprising  
CC a joined sequence of at least 24 nucleotides but not more than 70  
CC nucleotides and having secondary structure defined by: (a) 3 nucleotides  
CC forming a first side of a first double stranded (ds) region; (b) 2  
CC nucleotides forming a first side of an internal loop region; (c) 4  
CC nucleotides forming a first side of a second ds region; (d) 4 or 5  
CC nucleotides forming an end loop region; (e) 4 nucleotides forming a  
CC second side of the second ds region; (f) 4 nucleotides forming a second  
CC side of the internal loop region; and (g) 3 nucleotides forming a second  
CC side of the first ds region; (2) a purified and isolated RNA fragment  
CC comprising the human sequence UUUACACAUACUUGUUCAGAAAUUC (11). The  
CC methods and products can be used for identifying agents which modulate  
CC the activity of biomolecules, particularly RNA. Such agents can be used  
CC as pharmaceutical, agricultural or industrial compounds.  
XX  
SQ Sequence 42 BP; 11 A; 10 C; 7 G; 14 T; 0 other;  
  
Query Match 62.18; Score 18; DB 21; Length 42;  
Best Local Similarity 54.2%; Pred. No. 1;  
Matches 13; Conservative 5; Mismatches 6; Indels 0; Gaps 0;  
  
QY 4 gauncuununguagcccnangng 27  
||:|:::|:||||| | | |  
Db 7 gatcttcttgtaagcctcaggg 30  
  
RESULT 15  
AAA71126  
ID AAA71126 standard; RNA: 42 BP.  
XX  
AC AAA71126;  
XX  
DT 27-APR-2001 (first entry)  
XX  
DE Molecular interaction site RNA #195.  
XX  
KM Modulator; identification; molecular interaction; virtual library; ss.  
XX  
OS Unidentified.  
XX  
PN W09958947-A2.  
XX  
PD 18-NOV-1999.  
XX  
PE 12-MAY-1999; 99WO-US10361.  
XX  
PR 12-MAY-1998; 98US-0076404.  
XX  
PR 12-MAY-1998; 98US-0085092.  
XX  
PA (ISIS-) ISIS PHARM INC.  
XX  
PI Ecker DJ, Griffey R, Crooke ST, Sampath R, Swayze E, Mohan V;  
PI Hofstadler S, McNeil J;  
XX  
DR WPI; 2000-086439/07.  
XX  
PT Identifying compounds which modulate activity of target biomolecules,  
PT used to provide compounds which can be used as pharmacological.

PT agricultural and industrial compounds -  
XX  
PS Example 7; Figure 126; 405pp; English.

CC This invention describes a novel method for identifying compounds which  
CC modulate the activity of a target biomolecule. The method uses  
CC 3-dimensional representations of the biomolecule and a library of  
CC compounds and comprises (a) identifying at least one molecular  
CC interaction site of the target RNA; (b) generating in silico a virtual  
CC library of compounds predicted or calculated to interact with the  
CC molecular interaction site; and (c) comparing 3-dimensional (3-D)  
CC representations of the target RNA with members of the virtual library of  
CC compounds to generate a hierarchy of the compounds ranked in accordance  
CC with their respective ability to form physical interactions with the  
CC molecular interaction site. The method also describes (1) RNA comprising  
CC a joined sequence of at least 24 nucleotides but not more than 70  
CC nucleotides and having secondary structure defined by: (a) 3 nucleotides  
CC forming a first side of a first double stranded (ds) region; (b) 2  
CC nucleotides forming a first side of an internal loop region; (c) 4  
CC nucleotides forming a first side of a second ds region; (d) 4 or 5  
CC nucleotides forming an end loop region; (e) 4 nucleotides forming a  
CC second side of the second ds region; (f) 4 nucleotides forming a second  
CC side of the internal loop region; and (g) 3 nucleotides forming a second  
CC side of the first ds region; (2) a purified and isolated RNA fragment  
CC comprising the human sequence UUUACACUAUAUUCUGUUUACAGAAAUUC (11). The  
CC methods and products can be used for identifying agents which modulate  
CC the activity of biomolecules, particularly RNA. Such agents can be used  
CC as pharmaceutical, agricultural or industrial compounds.

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Query Match      62.1%; Score 18; DB 21; Length 42;
Best Local Similarity 75.0%; Pred. No. 1;
Matches 18; Conservative 0; Mismatches 6; Indels 0;
Gaps 0;

QY      4 gauncuununguaagcccnangng 27
      ||| |||| | ||||| | |
Db       7 gauncuununguaagcccnacgag 30

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Search completed: April 29, 2002, 22:45:09  
Job time: 4035 sec

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GenCore version 4.5  
Copyright (c) 1993 - 2000 CompuGen Ltd.

OM nucleic - nucleic search, using sw model

Run on: April 29, 2002, 19:50:29 ; Search time 60.62 Seconds  
(without alignments)  
117.509 Million cell updates/sec

Title: US-09-310-844C-23  
Perfect score: 29  
Sequence: 1 nngaucunungnaagccnangnn 29

Scoring table: IDENTITY NUC  
Gapop 10.0 , Gapext 1.0

Searched: 383533 seqs, 122816752 residues

Total number of hits satisfying chosen parameters: 613726

Minimum DB seq length: 0  
Maximum DB seq length: 100

Post-processing: Minimum Match 0%  
Maximum Match 100%  
Listing first 45 summaries

Database : Issued\_Patents\_NA:\*  
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2: /cgn2\_6/ptodata/1/ina/5B\_COMB.seq:\*  
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4: /cgn2\_6/ptodata/1/ina/6B\_COMB.seq:\*  
5: /cgn2\_6/ptodata/1/ina/6CTUS\_COMB.seq:\*  
6: /cgn2\_6/ptodata/1/ina/backfiles1.seq:\*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

## SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
1	12.2	42.1	27	6 5258283-10	Patent No. 5258283
2	12.2	42.1	69	2 US-08-410-654B-30	Sequence 30, Appl
3	12.2	42.1	69	2 US-08-474-851-30	Sequence 30, Appl
4	12.2	42.1	69	2 US-08-481-560-30	Sequence 30, Appl
5	11.8	40.7	21	2 US-08-747-536-10	Sequence 10, Appl
6	11.6	40.0	36	4 US-08-218-369-7	Sequence 7, Appl
7	11.6	40.0	36	4 US-08-218-369-15	Sequence 15, Appl
8	11.6	40.0	36	5 PCT-US95-03742-7	Sequence 7, Appl
9	11.6	40.0	36	5 PCT-US95-03742-15	Sequence 15, Appl
10	11.2	38.6	25	1 US-08-741-881-28	Sequence 28, Appl
11	11.2	38.6	25	1 US-08-739-158-28	Sequence 28, Appl
12	11.2	38.6	25	1 US-08-739-157-28	Sequence 28, Appl
13	11.2	38.6	25	3 US-08-404-796-28	Sequence 28, Appl
14	11.2	38.6	25	3 US-08-931-869-28	Sequence 28, Appl
15	11.2	38.6	25	3 US-09-350-399-28	Sequence 28, Appl
16	11.2	38.6	33	1 US-08-741-881-29	Sequence 29, Appl
17	11.2	38.6	33	1 US-08-739-158-29	Sequence 29, Appl
18	11.2	38.6	33	3 US-08-739-167-29	Sequence 29, Appl
19	11.2	38.6	33	3 US-08-404-796-29	Sequence 29, Appl
20	11.2	38.6	33	3 US-08-931-869-29	Sequence 29, Appl
21	11.2	38.6	33	3 US-09-350-399-29	Sequence 29, Appl
22	11.2	38.6	36	2 US-08-642-045B-17	Sequence 17, Appl
23	11.2	38.6	36	3 US-08-852-268-17	Sequence 17, Appl
24	11.2	38.6	70	4 US-09-364-380-29	Sequence 29, Appl
25	11.2	37.9	31	1 US-08-323-531-71	Sequence 71, Appl
26	11.2	37.9	31	1 US-08-198-094-71	Sequence 71, Appl
27	11.2	37.9	31	3 US-08-480-640A-119	Sequence 119, App

c 28	11	37.9	31	3	US-08-295-802-119	Sequence 119, App
c 29	11	37.9	31	4	US-08-107-794A-71	Sequence 71, Appl
c 30	11	37.9	31	4	US-08-488-237A-119	Sequence 119, App
c 31	11	37.9	31	4	US-08-375-992A-119	Sequence 119, App
c 32	11	37.9	31	5	PCT-US93-07424-71	Sequence 71, Appl
c 33	11	37.9	31	5	PCT-US95-02087-71	Sequence 71, Appl
c 34	11	37.9	70	2	US-08-488-402A-127	Sequence 127, App
c 35	11	37.9	70	2	US-08-484-552A-127	Sequence 127, App
c 36	11	37.9	70	5	PCT-US96-09472-127	Sequence 127, App
c 37	10.8	37.2	19	1	US-08-365-109B-1	Sequence 1, Appl
c 38	10.8	37.2	19	1	US-08-365-109B-3	Sequence 3, Appl
c 39	10.8	37.2	20	4	US-09-560-594-53	Sequence 53, Appl
c 40	10.8	37.2	25	4	US-08-943-731-336	Sequence 336, App
c 41	10.8	37.2	27	4	US-09-253-396A-137	Sequence 137, App
c 42	10.8	37.2	45	1	US-08-171-389-130	Sequence 130, App
c 43	10.8	37.2	45	1	US-08-171-389-342	Sequence 342, App
c 44	10.8	37.2	45	1	US-08-123-936-130	Sequence 130, App
c 45	10.8	37.2	45	1	US-08-123-936-342	Sequence 342, App

## ALIGNMENTS

RESULT 1  
5258283-10  
Patent No. 5258283  
APPLICANT: FRAZIER, MARVIN E.; MALAVIA, LOUIS P.; SAMUEL, JAMES E.; BACA, OSWALD G.  
TITLE OF INVENTION: DETECTION AND DIFFERENTIATION OF COXIELLA BURNETII IN BIOLOGICAL FLUIDS  
NUMBER OF SEQUENCES: 17  
CURRENT APPLICATION DATA:  
APPLICATION NUMBER: US/07/425, 856  
FILING DATE: 23-Oct-1999  
PRIOR APPLICATION DATA:  
APPLICATION NUMBER: 927, 779  
FILING DATE: 05-NOV-1986  
APPLICATION NUMBER: 795, 207  
FILING DATE: 05-NOV-1985  
SRQ ID NO:10:  
LENGTH: 27  
5258283-10

Query Match 42.1%; Score 12.2; DB 6; Length 27;  
Best Local Similarity 40.9%; Pred. No. 1.4e+02;  
Matches 9; Conservative 5; Mismatches 8; Indels 0; Gaps 0;  
QY 4 gauncunungnaagccnang 25  
DB 4 ggtcttgataagccaatg 25

RESULT 2  
US-08-410-654B-30  
Sequence 30, Application US/08410654B  
Patent No. 5833976  
GENERAL INFORMATION:  
APPLICANT: Rene de Waal Malefyt  
APPLICANT: Di-Hwei Hsu  
APPLICANT: Anne O'Garra  
APPLICANT: Hergen Splits  
TITLE OF INVENTION: Use of Interleukin-10 to Treat  
NUMBER OF SEQUENCES: 61  
CORRESPONDENCE ADDRESS:  
ADDRESSER: Schering-Plough Corporation  
STREET: 2000 Galloping Hill Road  
CITY: Kenilworth  
STATE: New Jersey  
COUNTRY: USA  
ZIP: 07033  
COMPUTER READABLE FORM:

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CEDUM TYPE: Floppy disk
COMPUTER: Macintosh
OPERATING SYSTEM: 7.5.3
SOFTWARE: Microsoft Word 5.1a
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/410,654B
FILING DATE: 24-MAR-1995
CLASSIFICATION: 424
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 08/229,854
FILING DATE: 19-APR-1994
APPLICATION NUMBER: US 07/926,853
FILING DATE: 06-AUG-1992
APPLICATION NUMBER: US 07/742,129
FILING DATE: 06-AUG-1991
ATTORNEY/AGENT INFORMATION:
NAME: Foukje, Cynthia L.
REGISTRATION NUMBER: 32,364
REFERENCE/DOCKET NUMBER: DX0221K01
TELECOMMUNICATION INFORMATION:
TELEPHONE: 908-298-7987
TELEFAX: 908-298-5388
INFORMATION FOR SEQ ID NO: 30:
SEQUENCE CHARACTERISTICS:
LENGTH: 69 base pairs
TYPE: nucleic acid
STRANDEDNESS: double
TOPOLOGY: linear
MOLECULE TYPE: DNA (oligonucleotide)
OS-08-410-654B-30

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Query Match	42.1%	Score 12.2;	DB 2;	Length 69;
Best Local Similarity	43.5%;	Pred. No. 1.7e+02;		
Matches 10; Conservative	4;	Mismatches 9;	Indels	

Qy 5 auncuunguaagcccnangng 27  
| : | : : | | | | | |  
Db 11 ATGCCTTAATAAGCTCCAAGAG 33

RESULT 3  
US-08-474-851-30  
Sequence 30, Application US/08474851  
Patient No. 5837232  
GENERAL INFORMATION:  
APPLICANT: Rene de Waal Malefyt  
APPLICANT: Di-Hwei Hsu  
APPLICANT: Anne O'Garra  
APPLICANT: Hergen Spits  
TITLE OF INVENTION: Use of An Interleukin-10 Antagonist to Treat  
TITLE OF INVENTION: A B Cell Mediated Autoimmune Disorder  
NUMBER OF SEQUENCES: 61  
CORRESPONDENCE ADDRESS:  
ADDRESSEE: Schering-Plough Corporation  
STREET: 2000 Galloping Hill Road  
CITY: Kenilworth  
STATE: New Jersey  
COUNTRY: USA  
ZIP: 07033  
COMPUTER READABLE FORM:  
MEDIUM TYPE: Floppy disk  
COMPUTER: Macintosh  
OPERATING SYSTEM: 7.5.3  
SOFTWARE: Microsoft Word 6.0  
CURRENT APPLICATION DATA:  
APPLICATION NUMBER: US/08/474,851  
FILING DATE: 07-JUN-1995  
CLASSIFICATION: 424  
PRIOR APPLICATION DATA:  
APPLICATION NUMBER: 08/410,654  
FILING DATE: 24-MAR-1995  
APPLICATION NUMBER: US 08/229,854

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1      FILING DATE: 19-APR-1994
2      APPLICATION NUMBER: US 07/926, 853
3      FILING DATE: 06-AUG-1992
4      APPLICATION NUMBER: US 07/742, 129
5      FILING DATE: 06-AUG-1991
6      ATTORNEY/AGENT INFORMATION:
7      NAME: Foulke, Cynthia L.
8      REGISTRATION NUMBER: 32,364
9      REFERENCE/DOCKET NUMBER: DX0221K01GD
10     TELECOMMUNICATION INFORMATION:
11     TELEPHONE: 908-298-2987
12     TELEFAX: 908-298-5388
13     INFORMATION FOR SEQ. ID. NO.: 30:
14     SEQUENCE CHARACTERISTICS:
15     LENGTH: 69 base pairs
16     TYPE: nucleic acid
17     STRANDEDNESS: double
18     TOPOLOGY: linear
19     MOLECULE TYPE: DNA (oligonucleotide)
20     US-08-474-851-30

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Query Match	42.1%;	Score 12.2;	DB 2;	Length 69;
Best Local Similarity	43.5%;	Pred. No. 1.7e+02;		
Matches 10; Conservative	4;	Mismatches 9;	Indels 0;	Gaps 0;

QY 5 auncuuunguaagcccnang 27  
| : | : : | | | | |  
Db 11 ATGCCTTTAATAAGCTCCAAGAG 33

RESULT 4  
US-08-481-560-30  
Sequence 30, Application US/08481560  
Patent No. 5637293  
GENERAL INFORMATION:  
APPLICANT: Rene de Waal Malefyt  
APPLICANT: Di-Hwei Hsu  
APPLICANT: Anne O'Garra  
APPLICANT: Hergen Spits  
TITLE OF INVENTION: Use of Interleukin-10 to Modulate  
TITLE OF INVENTION: Inflammation or T-cell Mediated  
TITLE OF INVENTION: Immune Function  
NUMBER OF SEQUENCES: 61  
CORRESPONDENCE ADDRESSES:  
ADDRESSEE: Schering-Plough Corporation  
STREET: 2000 Galloping Hill Road  
CITY: Kenilworth  
STATE: New Jersey  
COUNTRY: USA  
ZIP: 07033  
COMPUTER READABLE FORM:  
MEDIUM TYPE: Floppy disk  
COMPUTER: Macintosh  
OPERATING SYSTEM: 7.5.3  
SOFTWARE: Microsoft Word 6.0  
CURRENT APPLICATION DATA:  
APPLICATION NUMBER: US/08/481,560  
FILING DATE: 07-JUN-1995  
CLASSIFICATION: 424  
PRIOR APPLICATION DATA:  
APPLICATION NUMBER: 08/410,654  
FILING DATE: 24-MAR-1995  
APPLICATION NUMBER: US 08/229,854  
FILING DATE: 19-APR-1994  
APPLICATION NUMBER: US 07/926,853  
FILING DATE: 06-AUG-1992  
APPLICATION NUMBER: US 07/742,129  
FILING DATE: 06-AUG-1991  
ATTORNEY/AGENT INFORMATION:  
NAME: Foulke, Cynthia L.  
REGISTRATION NUMBER: 32,364  
REFERENCE/DOCKET NUMBER: DX0221KOIGC



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STREET: 1100 Peachtree Street, Suite 2800
CITY: Atlanta
STATE: Georgia
COUNTRY: USA
ZIP: 30309-4530
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patent Release #1.0, Version #1.25
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/218,369
FILING DATE: 28-MAR-1994
CLASSIFICATION: 435
ATTORNEY/AGENT INFORMATION:
NAME: Pabst, Patrea L.
REGISTRATION NUMBER: 31,284
TELECOMMUNICATION INFORMATION:
TELEPHONE: (404) 815-6508
TELEFAX: (404) 815-6555
INFORMATION FOR SEQ ID NO: 15:
SEQUENCE CHARACTERISTICS:
LENGTH: 36 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
MOLECULE TYPE: DNA
HYPOTHETICAL: NO
ANTI-SENSE: NO
FEATURE:
NAME/KEY: misc_feature
LOCATION: 1..36
OTHER INFORMATION: /note= "Nucleotides 5 through 36 are complementary to nucl
US-08-218-369-15

Query Match          40.0%; Score 11.6; DB 4; Length 36;
Best Local Similarity 41.7%; Pred. No. 3.4e+02;
Matches 10; Conservative 4; Mismatches 10; Indels 0; Gaps 0;

QY 4 gauncuununguaagcccnang 27
|||:::|:|11111111
Db 10 GAAGCTTAGTGCGGCCCATGAG 33

RESULT 8
-US95-03742-7/C
Sequence 7, Application PC/TUS9503742
GENERAL INFORMATION:
APPLICANT: The UAB Research Foundation
TITLE OF INVENTION: Ligands Added to Adenovirus Fiber
NUMBER OF SEQUENCES: 18
CORRESPONDENCE ADDRESS:
ADDRESSER: Patrea L. Pabst
STREET: 2800 One Atlantic Center
STREET: 1201 West Peachtree Street
CITY: Atlanta
STATE: Georgia
COUNTRY: USA
ZIP: 30309-3450
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patent Release #1.0, Version #1.25
CURRENT APPLICATION DATA:
APPLICATION NUMBER: PCT/US95/03742
FILING DATE:
CLASSIFICATION:
ATTORNEY/AGENT INFORMATION:
NAME: Pabst, Patrea L.
REGISTRATION NUMBER: 31,284
TELECOMMUNICATION INFORMATION:
TELEPHONE: (404) 873-8795
TELEFAX: (404) 873-8795
INFORMATION FOR SEQ ID NO: 15:
SEQUENCE CHARACTERISTICS:
LENGTH: 36 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
MOLECULE TYPE: DNA
HYPOTHETICAL: NO
ANTI-SENSE: NO
FEATURE:
NAME/KEY: misc_feature
LOCATION: 1..36
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REFERENCE/DOCKET NUMBER: IG1101
TELECOMMUNICATION INFORMATION:
TELEPHONE: (404) 873-8794
TELEFAX: (404) 873-8795
INFORMATION FOR SEQ ID NO: 7:
SEQUENCE CHARACTERISTICS:
LENGTH: 36 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
MOLECULE TYPE: DNA
HYPOTHETICAL: NO
ANTI-SENSE: NO
FEATURE:
NAME/KEY: misc_feature
LOCATION: 1..36
OTHER INFORMATION: /note= "Nucleotide sequence
PCT-US95-03742-7

Query Match          40.0%; Score 11.6; DB 5; Length 36;
Best Local Similarity 41.7%; Pred. No. 3.4e+02;
Matches 10; Conservative 4; Mismatches 10; Indels 0; Gaps 0;

QY 4 gauncuununguaagcccnang 27
|||:::|:|11111111
Db 31 GAAGCTTAGTGCGGCCCATGAG 8

RESULT 9
PCT-US95-03742-15
Sequence 15, Application PC/TUS9503742
GENERAL INFORMATION:
APPLICANT: The UAB Research Foundation
TITLE OF INVENTION: Ligands Added to Adenovirus Fiber
NUMBER OF SEQUENCES: 18
CORRESPONDENCE ADDRESS:
ADDRESSER: Patrea L. Pabst
STREET: 2800 One Atlantic Center
STREET: 1201 West Peachtree Street
CITY: Atlanta
STATE: Georgia
COUNTRY: USA
ZIP: 30309-3450
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patent Release #1.0, Version #1.25
CURRENT APPLICATION DATA:
APPLICATION NUMBER: PCT/US95/03742
FILING DATE:
CLASSIFICATION:
ATTORNEY/AGENT INFORMATION:
NAME: Pabst, Patrea L.
REGISTRATION NUMBER: 31,284
REFERENCE/DOCKET NUMBER: IG1101
TELECOMMUNICATION INFORMATION:
TELEPHONE: (404) 873-8794
TELEFAX: (404) 873-8795
INFORMATION FOR SEQ ID NO: 15:
SEQUENCE CHARACTERISTICS:
LENGTH: 36 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
MOLECULE TYPE: DNA
HYPOTHETICAL: NO
ANTI-SENSE: NO
FEATURE:
NAME/KEY: misc_feature
LOCATION: 1..36
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OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patent Release #1.0, Version #1.25
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/739,167
FILING DATE: 30-OCT-1996
CLASSIFICATION: 435
ATTORNEY/AGENT INFORMATION:
NAME: Mcmasters, David D.
REGISTRATION NUMBER: 33,963
REFERENCE/DOCKET NUMBER: 930049,423C7 / 1146,0008
TELECOMMUNICATION INFORMATION:
TELEPHONE: (206) 622-4500
TELEFAX: (206) 682-6031
INFORMATION FOR SEQ ID NO: 28:
SEQUENCE CHARACTERISTICS:
LENGTH: 25 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear

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Query Match	38.6%	Score 11.2	DB 2	Length 25
Best Local Similarity	36.4%	Pred. No.	5.4e+02	
Matches	8	Conservative	5	Mismatches 9
				Indels 0
				Gaps 0

Qy 6 uncuuunguaagcccnangng 27  
: |:: |: ||| |  
Db 24 TCCTTTAGGTTAGCCGTACAAG 3

RESULT 13  
US-08-404-796-28/C  
; Sequence 28, Application US/08404796

GENERAL INFORMATION:  
 APPLICANT: Dubensky Jr, Thomas W  
 APPLICANT: Polo, John M.  
 APPLICANT: Ibanez, Carlos E.  
 APPLICANT: Chaney, Stephen M.W.  
 APPLICANT: Jolly, Douglas J.  
 APPLICANT: Driver, David A.  
 APPLICANT: Belli, Barbara A.  
 TITLE OF INVENTION: EUKARYOTIC LAYERED VECTOR INITIATION SYSTEMS  
 NUMBER OF SEQUENCES: 128

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;      TOPOLOGY:  linear
US-08-404-796-28

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Query Match	38.6%	Pred 11.2;	DB 3;	Length 25;
Best Local Similarity	36.4%	Pred No. 5.4e+02;		
Matches	8;	Conservative	5;	Mismatches 9;
				Indels 0;
				Gaps 0;

QY 6 uncuuunguaagcccnangng 27  
: |: : |: |: |  
Db 24 TCCTTTAGGTTAGCCGTACAAG 3

RESULT 14  
US-08-931-869-28/c  
; Sequence 28, Application US/08931869  
; Patent No. 6015694

1 APPLICANT: Dubensky Jr, Thomas W  
 2  
 3 APPLICANT: Polo, John M.  
 4  
 5 APPLICANT: Idanez, Carlos E.  
 6  
 7 APPLICANT: Chang, Stephen M.W.  
 8  
 9 APPLICANT: Jolly, Douglas J.  
 10  
 11 APPLICANT: Driver, David A.  
 12  
 13 APPLICANT: Belli, Barbara A.  
 14  
 15 TITLE OF INVENTION: EUKARYOTIC LAYERED VECTOR INITIATION SYSTEMS  
 16  
 17 NUMBER OF SEQUENCES: 128  
 18  
 19 CORRESPONDENCE ADDRESS:

Query Match	38.6%	Score	11.2	DB	3	Length	25
Best Local Similarity	36.4%	Pred. No.	5.4e+02				
Matches	8	Conservative	5	Mismatches	9	Indels	0
						Gaps	0

QY 6 uncuuunguaagcccnangng 27  
: |:: |: ||| |  
Db 24 TCCTTTAGGTTAGCCGTACAAG 3

RESULT 15  
US-09-350-399-28/c

; Sequence 28, Application US/09350399  
; Patent No. 6342372  
; GENERAL INFORMATION:  
; APPLICANT: Dubensky Jr, Thomas W  
; Polo, John M.  
; Jolly, Douglas J.  
; Driver, David A.  
; TITLE OF INVENTION: EUKARYOTIC LAYERED VECTOR INITIATION SYSTEMS  
; NUMBER OF SEQUENCES: 128  
; CORRESPONDENCE ADDRESS:  
; ADDRESSEE: SEED and BERRY LLP  
; STREET: 6300 Columbia Center, 701 Fifth Avenue  
; CITY: Seattle  
; STATE: Washington  
; COUNTRY: US  
; ZIP: 98104-7092  
; COMPUTER READABLE FORM:  
; MEDIUM TYPE: Floppy disk  
; COMPUTER: IBM PC compatible  
; OPERATING SYSTEM: PC-DOS/MS-DOS  
; SOFTWARE: PatentIn Release #1.0, Version #1.25  
; CURRENT APPLICATION DATA:  
; APPLICATION NUMBER: US/09/350,399  
; FILING DATE: 08-Jul-1999  
; CLASSIFICATION: <Unknown>  
; ATTORNEY/AGENT INFORMATION:  
; NAME: McMasters, David D.  
; REGISTRATION NUMBER: 33,963  
; REFERENCE/DOCKET NUMBER: 930049.423D1 / 1146.010  
; TELECOMMUNICATION INFORMATION:  
; TELEPHONE: (206) 622-4900  
; TELEFAX: (206) 682-6031  
; INFORMATION FOR SEQ ID NO: 28:  
; SEQUENCE CHARACTERISTICS:  
; LENGTH: 25 base pairs  
; TYPE: nucleic acid  
; STRANDEDNESS: single  
; TOPOLOGY: linear  
; SEQUENCE DESCRIPTION: SEQ ID NO: 28:  
; US-09-350-399-28

Query Match 38.6%; Score 11.2; DB 4; Length 25;  
Best Local Similarity 36.4%; Pred. No. 5.4e+02;  
Matches 8; Conservative 5; Mismatches 9; Indels 0; Gaps 0;  
QY 6 uncuununguagcccnangng 27  
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Db 24 TCCTTAGTGTAGCCGTACAAG 3

Search completed: April 29, 2002, 22:39:48  
Job time: 10159 sec

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GenCore version 4.5  
Copyright (c) 1993 - 2000 Compugen Ltd.

OM nucleic - nucleic search, using sw model

Run on: April 29, 2002, 18:34:27 ; Search time 1874.63 Seconds  
(without alignments)  
208.794 Million cell updates/sec

Title: US-09-310-844C-23  
Perfect score: 29  
Sequence: 1 nngauuncununguaagccnangnngn 29

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Searched: 13736307 seqs, 674847542 residues

Total number of hits satisfying chosen parameters: 297742

Minimum DB seq length: 0  
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Maximum Match 100%  
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9: gb\_est1:\*  
10: gb\_est2:\*  
11: gb\_hic:\*  
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13: em\_gss\_hum:\*  
14: em\_gss\_inv:\*  
15: em\_gss\_pln:\*  
16: em\_gss\_vrt:\*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

## SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	13	44.8	46	12	AZ833686 2M0115L20
2	12.8	44.1	76	12	A0025263 EP(3)3081
3	12.8	44.1	86	12	CNS02IOD
4	12.4	42.8	52	9	AA700959
5	12.4	42.8	70	9	AA468615
6	12.2	42.1	48	12	A2503560
7	12.2	42.1	75	12	A2453746
8	12.2	42.1	86	10	C01512
9	12.2	42.1	92	10	B1305219
10	12.2	42.1	96	10	H81976
11	12.2	42.1	98	12	A2767813
12	11.8	40.7	40	9	AA975071
13	11.8	40.7	49	10	BE970036
14	11.8	40.7	65	9	AA733449
15	11.8	40.7	70	9	A1767928
16	11.8	40.7	70	12	BH216023
17	11.8	40.7	75	12	CNS01561

C 18	11.8	40.7	83	10	BE845147	BE845147 AD07E1277
C 19	11.8	40.7	86	10	BE845145	BE845145 AD07E06T7
C 20	11.8	40.7	87	12	AA807716	AA807716 2M0070D13
C 21	11.8	40.7	93	9	AA488835	AA488835 aa54h10.r
C 22	11.8	40.7	95	12	A2598556	A2598556 1M0413K23
C 23	11.8	40.7	98	9	AW311302	AW311302 sg35b11.y
C 24	11.6	40.0	49	10	U44334	U44334 END044334 As
C 25	11.6	40.0	58	9	A1584456	A1584456 fb93h12.x
C 26	11.6	40.0	70	9	A1E14489	A1E14489 wj73g11.x
C 27	11.6	40.0	79	12	A2336769	A2336769 1M0067L12
C 28	11.6	40.0	81	12	BH627770	BH627770 1007076E0
C 29	11.6	40.0	85	9	AA617776	AA617776 np99e08.s
C 30	11.6	40.0	85	10	W20254	W20254 zb42a10.r1
C 31	11.6	40.0	93	12	A2832088	A2832088 2M0112D15
C 32	11.6	40.0	94	9	AA558052	AA558052 n118f06.s
C 33	11.6	40.0	97	9	A1181481	A1181481 uc62g11.r
C 34	11.6	40.0	100	9	A1148625	A1148625 qc62g05.x
C 35	11.6	40.0	100	10	H13996	H13996 EST00022 Ch
C 36	11.4	39.3	32	12	HSMC42B09	X88068 H.sapiens D
C 37	11.4	39.3	40	12	TA253H010	AT483109 T. brucei
C 38	11.4	39.3	46	9	A1867082	A1867082 w196e09.x
C 39	11.4	39.3	66	9	AA247859	AA247859 j3371.seq
C 40	11.4	39.3	67	12	TA113E04Q	AL460350 T. brucei
C 41	11.4	39.3	73	9	A1900474	A1900474 sc11b10.y
C 42	11.4	39.3	75	12	BH609927	BH609927 HIV22C11
C 43	11.4	39.3	76	9	AA988261	AA988261 os16a07.s
C 44	11.4	39.3	81	9	AA738756	AA738756 vv67B08.r
C 45	11.4	39.3	85	12	A2830379	A2830379 2M0109A09

## ALIGNMENTS

RESULT 1  
A2833686 46 bp DNA linear GSS 20-FEB-2001  
LOCUS 2M0115L20R Mouse 10kb plasmid UGCC1M library Mus musculus genomic  
DEFINITION clone UGCC2M0115L20 R, DNA sequence.

ACCESSION A2833686  
VERSION A2833686.1 GI:13003594  
KEYWORDS GSS.

SOURCE house mouse.  
ORGANISM Mus musculus

REFERENCE Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.  
AUTHORS Dunn,D., Aoyagi,A., Barber,M., Beacorn,T., Duval,B., Hamil,C., Islam,H., Longacre,S., Mahmoud,M., Meenen,E., Pedersen,T., Reilly,M., Rose,M., Rose,R., Stokes,R., Tingey,A., von Niederhausern,A. and Wright,D., Weiss,R.

TITLE Mouse whole genome scaffolding with paired end reads from 10kb plasmid inserts  
JOURNAL Unpublished (2000)  
COMMENT Contact: Robert B. Weiss  
University of Utah Genome Center  
University of Utah  
Rm. 308, Biomedical Polymers Research Bldg., 20 S. 2030 E., SLIC, UT  
84112, USA  
Tel: 801 585 5606  
Fax: 801 585 7177  
Email: ddunn@genetics.utah.edu  
Insert length: 10000 Std Error: 0.00  
Plate: 0115 row: L column: 20  
Seq primer: CACACGAGAAACGCTATGACC  
Class: plasmid ends  
High quality sequence stop: 46.  
Location/Qualifiers  
1. 46

FEATURES  
source  
/organism="Mus musculus"  
/strain="C57BL/6J"  
/db\_xref="taxon:10090"  
/clone="UGCC2M0115L20"  
/clone\_lib="Mouse 10kb plasmid library"

/sex="Male"  
/lab\_host="E. Coli strain XL10-Gold, T1-resistant, F-"  
/note="Vector: pMD42nv; Purified genomic DNA from M.  
musculus C57BL/6J (male) was obtained from the Jackson  
Laboratory Mouse DNA Resource  
(http://www.jax.org/resources/documents/dnares/). The DNA  
was hydrodynamically sheared by repeated passage through a  
0.005 inch orifice at constant velocity. The sheared DNA  
was blunt end-repaired with T4 DNA polymerase and T4  
polynucleotide kinase. Adaptor oligonucleotides were  
ligated to the blunt ends in high molar excess. The  
adapted DNA was purified and size-selected for a 9.5 to  
10.5 kb range using preparative agarose gel  
electrophoresis. Vector DNA was prepared from a derivative  
of pMD42 (g1473214|gb|AF129072.1), a copy-number  
inducible derivative of plasmid R1. The vector was ligated  
with adaptors complementary to the insert adaptors and  
purified. The sheared, adapted mouse DNA was annealed to  
chemically-competent E. coli XL10-Gold (Stratagene) cells  
and selected for ampicillin resistance."

BASE COUNT  
ORIGIN  
14 a 6 c 13 g 13 t

Query Match 44.8%; Score 13; DB 12; Length 46;  
Best Local Similarity 50.0%; Pred. No. 1.4e+04;  
Matches 8; Conservative 5; Mismatches 3; Indels 0; Gaps 0;

QY 4 gauncununguaac 19  
||:|:::|:||||  
Db 8 GATACCTTAAGTAAAGC 23

RESULT 2  
A0025263/c 76 bp DNA linear GSS 23-AUG-2000  
DEFINITION  
A0025263 Drosophila melanogaster EP line Drosophila melanogaster  
genomic Sequence recovered from 5' end of P element, DNA sequence.  
ACCESSION  
A0025263  
VERSION  
A0025263.1 GI:3265615  
KEYWORDS  
GSS.  
SOURCE  
ORGANISM  
fruit fly.  
Drosophila melanogaster  
Eukaryota; Metazoa; Arthropoda; Tracheata; Hexapoda; Insecta;  
Pterygota; Neoptera; Endopterygota; Diptera; Brachycera;  
Muscomorpha; Ephydroidea; Drosophilidae; Drosophila.  
1 (bases 1 to 76)  
Liao, G.-C., Rehm, E.J. and Rubin, G.M.  
Insertion site preferences of the P transposable element in  
Drosophila melanogaster  
Proc. Natl. Acad. Sci. U.S.A. 97 (7), 3347-3351 (2000)  
20202638  
Contact: Gerald Rubin  
Berkeley Drosophila Genome Project  
University of California, Berkeley  
LSA Building, Berkeley, CA 94720-3200, USA  
Fax: 5106433947  
Email: gerry@fruitfly.berkeley.edu  
Sequence recovery method was inverse PCR.  
Sequence orientation is forward strand relative to 5' end of P  
element

The P element insertion position is base 69 in the 76 bases. This  
insertion position refers to the first base of the 8 base target  
recognition sequence.  
Class: transposon-tagged.  
Location/Qualifiers  
1..76  
/organism="Drosophila melanogaster"  
/db\_xref="taxon:7227"  
/clone\_id="Drosophila melanogaster EP line"

FEATURES  
SOURCE

1..76  
/organism="Drosophila melanogaster"  
/db\_xref="taxon:7227"  
/clone\_id="Drosophila melanogaster EP line"

/note="Inverse PCR was performed on Drosophila  
melanogaster strains each of which contains a single EP  
transposable element insertion. (The generation of these  
insertion strains is described in Roth P, Szabo K, Bailey  
A, Laverly T, Rehm J, Rubin GM, Weigmann K, Milan M, Benes  
V, Ansoerg W, Cohen SM. 1998. Systematic gain-of-function  
genetics in Drosophila. Development 6:1049-1057.) The  
resultant fragment for each strain was directly sequenced  
to determine the genomic sequence at the site of  
insertion. Details of the protocols used can be found at  
http://fruitfly.berkeley.edu/P\_disrupt/inverse\_pcr.html."

BASE COUNT  
ORIGIN  
31 a 11 c 13 g 21 t

Query Match 44.1%; Score 12.8; DB 12; Length 76;  
Best Local Similarity 42.9%; Pred. No. 1.8e+04;  
Matches 9; Conservative 5; Mismatches 7; Indels 0; Gaps 0;

QY 5 auncununguaagccnang 25  
||:|:::|:||||  
Db 62 ATACTTATTTAATCCCAAG 42

RESULT 3  
CNS0210D 86 bp DNA linear GSS 13-MAY-2000  
LOCUS  
DEFINITION  
Tetradon nigroviridis genome survey sequence T7 end of clone  
141P20 of library G from Tetradon nigroviridis, genomic survey  
sequence.

ACCESSION  
AL199174.1 GI:7837325  
VERSION  
GSS: genome survey sequence.  
KEYWORDS  
Tetradon nigroviridis.  
SOURCE  
ORGANISM  
Tetradon nigroviridis  
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
Actinopterygii; Neopterygii; Teleostei; Euteleostei; Neoteleostei;  
Acanthomorpha; Acanthopterygii; Percomorpha; Tetraodontiformes;  
Tetraodontidae; Tetraodon.

REFERENCE  
1 (bases 1 to 86)  
Roest-Crollius, H., Jallion, O., Dasilva, C., Fitzames, C., Fisher, C.,  
Bonneau, L., Billault, A., Quetier, F., Saurin, W., Bernot, A. and  
Weissenbach, J.  
Characterization and repeat analysis of the compact genome of the  
freshwater pufferfish Tetraodon nigroviridis

TITLE  
JOURNAL  
REFERENCE  
2 (bases 1 to 86)  
Roest-Crollius, H., Jallion, O., Dasilva, C., Bonneau, L., Fisher, C.,  
Bernot, A., Fitzames, C., Wincker, P., Brottier, P., Quetier, F.,  
Saurin, W. and Weissenbach, J.  
Human gene number estimate provided by genome wide analysis using  
Tetradon nigroviridis DNA sequence

UNPUBLISHED  
3 (bases 1 to 86)  
Genoscope.  
Direct Submission  
Submitted (12-APR-2000) to the EMBL/Genbank/DBJ databases  
This sequence is a single read and was generated as part of a large  
scale clone and sequencing project of the Tetradon nigroviridis  
genome. For more information, please take a look at  
http://www.genoscope.cns.fr/Tetraodon.

FEATURES  
SOURCE  
1..86  
/organism="Tetradon nigroviridis"  
/db\_xref="taxon:99883"  
/clone="141P20"  
/clone\_id="G"  
/note="Genoscope sequence ID : COAG141DH10UP1-end : T7"

BASE COUNT  
ORIGIN  
12 a 12 c 26 g 34 t 2 others

Query Match 44.1%; Score 12.8; DB 12; Length 86;



REFERENCE  
Mammalia; Eutheria; Primates; Catarrhini; Hominoidea; Homo.  
1 (bases 1 to 86)





Best Local Similarity	50.0%	Pred. No. 3.8e+04	
Matches 11: Conservative	3:	Mismatches 8:	Indels 0:
Gaps	0:		

RESULT	11
AZ767813/C	
LOCUS	
DEFINITION	98 bp DNA linear GSS 16-FEB-2001
ACCESSION	M0567G2AF Mouse 10kb plasmid UUC1M library Mus musculus genomic clone UUC1M0567G2A F, DNA sequence.
	AZ767813

REFERENCE	ORGANISM	SOURCE	KEYWORDS	VERSION	ACCESSION
Dunn, D., Aoyagi, A., Barber, M., Beacorn, T., Duval, B., Hamill, C.,	Mammalia, Eutheria, Rodentia, Sciurognathi; Muridae; Murinae; Mus	house mouse, Mus musculus		A2767813.1	GI:12886296

TITLE	Mouse whole genome scaffolding with paired end reads from 10kb
JOURNAL	plasmid inserts
COMMENT	Unpublished (2000)
	Contact: Robert B. Weiss

FEATURES	Location/Qualifiers
Source	1 00

		42.1%;	Score 12.2;	DB 12;	Length 98;
Query Match			Pred. No. 3.8e+04;		
Best Local Similarity		47.8%;			
Matches 11; Conservative 3; Mismatches 9; Indels 0; Gaps 0;					
OY	5	auncuuunnguagcccnangng	27		
	:	: :			
	73	ATCCCTACAGTAAGACTTAAAGGGG	51		
Db					

RESULT 12					
AA975071/c					
LOCUS	AA975071	40 bp	mRNA	linear	EST:26-AUG-1998
DEFINITION	0n03d07.s1 NCI-GCAP Kid3 Homo sapiens cDNA clone IMAGE:155597	3'			
	similar to TR:P70566 P70566 N-TROPOMODULIN.; mRNA sequence.				
ACCESSION	U00687				

ACCESSION VERSION KEYWORDS SOURCE ORGANISM	AA975071 AA975071 EST. human. Homo sapiens Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.	GI:3150863
REFERENCE	1 (bases 1 to 40)	

Trace considered overall poor quality  
Insert length: 2096 Std Error: 0.00  
Seq primer: -40m13 fwd. ET from Amersham  
High quality sequence stop: 1.

BASE COUNT	10 a	7 c	11 g	12 t
ORIGIN				
Query Match		40.7%;	Score 11.8;	DB 9;
Best Local Similarity	40.0%;	Pred. No. 5.8e+04;		Length 40;
Matches	8; Conservative	5; Mismatches	7; Indels	0; Gaps
QY	6 uncuununguaagcccnang	25		
	: : : : :   :			
Db	35 TCCTTCGCGTAAGACCTTGG	16		

BASE COUNT	17 a	32 c	22 g	27 t
ORIGIN				
<hr/>				
RESULT	13			
BE970036/c				
LOCUS	BE970036			
<hr/>				
49 bp	mRNA			
	linear			
	EST 04-OCT-2000			

DEFINITION 601680150F1 NIH\_MGC\_78 Homo sapiens cDNA clone IMAGE:3950172 5', mRNA sequence.

ACCESSION BE970036

VERSION BE970036.1 GI:10582969

KEYWORDS EST.

SOURCE human.

ORGANISM Homo sapiens

REFERENCE Mammalia; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; 1 (bases 1 to 49)

AUTHORS NIH-MGC <http://mhc.nci.nih.gov/>.

TITLE National Institutes of Health, Mammalian Gene Collection (MGC)

JOURNAL Unpublished (1999)

COMMENT Contact: Robert Strausberg, Ph.D.  
Email: [cgapbs-remail.nih.gov](mailto:cgapbs-remail.nih.gov)  
Tissue Procurement: CLONTECH Laboratories, Inc.  
cDNA Library Preparation: CLONTECH Laboratories, Inc.  
DNA Sequencing by: Incyte Genomics, Inc.  
Clone distribution: MGC clone distribution information can be found through the I.M.A.G.E. Consortium/LLNL at: <http://image.llnl.gov>  
Place: LBL816 row: d column: 13  
High quality sequence stop: 49.

FEATURES  
source  
1. 49  
Location/Qualifiers  
/organism="Homo sapiens"  
/db\_xref="taxon:9606"  
/clone="IMAGE:3950172"  
/clone\_11b="NIH\_MGC\_78"  
/lab\_host="DH10B (T1 Phage-resistant)"  
/note="Organ: pancreas; Vector: pDNR-LIB (Clontech); Site\_1: SfiI (ggcgccgcgcgc); Site\_2: SfiI (ggcgattatggc ); 5' and 3' adaptors were used in cloning as follows: 5' adaptor sequence: 5'-ATCTGAGAGCGGCGCGCGCATG-3' and 3' adaptor sequence: 5'-CACGCCATTATGACC-3' (where B = A, C, or G and N = A, C, G, or T). Average insert size 1.2 kb (range 0.5-4.0 kb). 14/15 colonies contained inserts by PCR. This library was enriched for full-length clones and was constructed by Clontech Laboratories (Palo Alto, CA)."

BASE COUNT 20 a 7 c 12 g 10 t

ORIGIN

Query Match 40.7%; Score 11.8; DB 10; Length 49;  
Best Local Similarity 45.0%; Pred. No. 5.9e+04;  
Matches 9; Conservative 4; Mismatches 7; Indels 0; Gaps 0;

6 uncuununguagccnang 25  
1 : : : : | | | | | | | |  
48 TTTTTCATGCAAGCCCCAGG 29

RESULT 14  
AA733449 65 bp mRNA linear EST 07-JAN-1998  
LOCUS AA733449/c  
DEFINITION vt73h08.r1 Barstead mouse irradiated colon MFLRB7 Mus musculus cDNA clone IMAGE:1176831 5' similar to gb:X06617 40S RIBOSOMAL PROTEIN S11 (HUMAN); mRNA sequence.

ACCESSION AA733449

VERSION AA733449.1 GI:2755116

KEYWORDS EST.

SOURCE house mouse.

ORGANISM Mus musculus

REFERENCE Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus. 1 (bases 1 to 65)

AUTHORS Marra, M., Hillier, L., Allen, M., Bowles, M., Dietrich, N., Dubaque, T., Geisel, S., Kucaba, T., Lacy, M., Le, M., Martin, J., Morris, M., Schellenberg, K., Steptoe, M., Tan, F., Underwood, K., Moore, B., Theising, B., Wylie, T., Lennon, G., Soares, B., Wilson, R. and Waterston, R.

TITLE The WashU-HIMI Mouse EST Project

JOURNAL Unpublished (1996)

COMMENT Contact: Marra M/Mouse EST Project  
WashU-HIMI Mouse EST Project  
Washington University School of Medicine  
4444 Forest Park Parkway, Box 8501, St. Louis, MO 63108  
Tel: 314 286 1800  
Fax: 314 286 1810  
Email: [mouseest@watson.wustl.edu](mailto:mouseest@watson.wustl.edu)  
This clone is available royalty-free through LLNL; contact the IMAGE Consortium ([info@image.llnl.gov](mailto:info@image.llnl.gov)) for further information.  
MGI:634679  
Trace considered overall poor quality  
Seq primer: -28m13 rev2 ET from Amersham  
High quality sequence stop: 1.

FEATURES  
source  
1. 65  
Location/Qualifiers  
/organism="Mus musculus"  
/strain="FVB/N"  
/db\_xref="taxon:10090"  
/clone="IMAGE:1176831"  
/clone\_11b="Barstead mouse irradiated colon MFLRB7"  
/dev\_stage="8 weeks"  
/lab\_host="DH10B"  
/note="Vector: pT73D-Pac (Pharmacia) with a modified polylinker. Site\_1: EcoRI; Site\_2: NotI. Tissue obtained from 8 week old mouse. Colon was harvested 72 hours after irradiation with 1400 Gys. 1st strand cDNA was primed with a Not I - 01190(dT) primer  
5' TGTTGCAATCTGAGTGGAGCGCGCGCCCTTTTTTTTTTTTTTTTTTTT  
T 3'; double-stranded cDNA was ligated to Eco RI adaptors (AATTCGATCCTTG), digested with Not I and cloned into the Not I and Eco RI sites of the modified pT773 vector. Library constructed by Bob Barstead."

BASE COUNT 23 a 15 c 16 g 11 t

ORIGIN

Query Match 40.7%; Score 11.8; DB 9; Length 65;  
Best Local Similarity 44.4%; Pred. No. 6e+04;  
Matches 8; Conservative 5; Mismatches 5; Indels 0; Gaps 0;

4 gauncuununguagccc 21  
1 : : : : | | | | | | | |  
44 GCTGCTTTTGCTAGACAC 27

RESULT 15  
A1767928 70 bp mRNA linear EST 21-DEC-1999  
LOCUS A1767928  
DEFINITION w199c01.x1 NCI-CGAP\_K1d12 Homo sapiens cDNA clone IMAGE:2401440 3' similar to SW:RT14\_HUMAN P78537 RT14 PROTEIN; mRNA sequence.

ACCESSION A1767928

VERSION A1767928.1 GI:5234426

KEYWORDS EST.

SOURCE human.

ORGANISM Homo sapiens

REFERENCE Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Homiidae; Homo. 1 (bases 1 to 70)

AUTHORS NCI-CGAP <http://www.ncbi.nlm.nih.gov/ncicgap>.

TITLE National Cancer Institute, Cancer Genome Anatomy Project (CGAP), Tumor Gene Index

JOURNAL Unpublished (1997)

COMMENT Contact: Robert Strausberg, Ph.D.  
Email: [cgapbs-remail.nih.gov](mailto:cgapbs-remail.nih.gov)  
Tissue Procurement: Christopher Moskaluk, M.D., Ph.D., Michael R. Emmert-Buck, M.D., Ph.D.  
cDNA Library Preparation: M. Bento Soares, Ph.D.  
DNA Sequencing by: Greg Lennon, Ph.D.  
Clone distribution: NCI-CGAP clone distribution information can be found through the I.M.A.G.E. Consortium/LLNL at:

www.bio.lnl.gov/db/rp/image/image.html

Trace considered overall poor quality  
Insert Length: 574 Std Error: 0.00  
Seq primer: -40UP from Gibco  
High quality sequence stop: 1.  
Location/Qualifiers

## FEATURES

Source

1. 70

/organism="Homo sapiens"

/db\_xref="taxon:9606"

/clone="IMAGE:2401440"

/clone\_lib="NCI\_CGAP\_Kid12"

/tissue\_type="2\_pooled tumors (clear cell type)"

/lab\_host="DH10B"

/note="Organ: kidney; Vector: p1773D-Pac (Pharmacia) with a modified polylinker; Site.1: Not I; Site.2: Eco RI; Plasmid DNA from the normalized library NCI\_CGAP\_Kid5 was prepared, and ss circles were made in vitro. Following HAP purification, this DNA was used as tracer in a subtractive hybridization reaction. The driver was PCR-amplified cDNAs from a pool of 5,000 clones made from the same library (clonoids 1323912-1325831, 1471368-1472903 and 1492104-1493255). Subtraction by Bento Soares and M. Fatima Bonaldo."

BASE COUNT 20 a 24 c 6 g 20 t  
ORIGIN

## Query Match

40.7%; Score 11.8; DB 9; Length 70;

Best Local Similarity 47.4%; Pred. No. 6e+04; 6; Indels 0; Gaps 0;  
Matches 9; Conservative 4; Mismatches 0;

5 auncununguaagccca 23  
|:|::| | | | | | |  
b 46 ATTCTTAAAGCAAGCCAGA 64

Search completed: April 29, 2002, 22:38:40  
Job time: 14653 sec